

S/188/62/000/004/010/010  
B108/B102

AUTHORS: Vasil'yev, S. S., Romanovskiy, Ye. A., Timushev, G. F.

TITLE: Cross section of 6.6-Mev proton absorption by  $F^{19}$  nuclei

PERIODICAL: Moscow. Universitet. Vestnik. Seriya III. Fizika, astronomiya, no. 4, 1962, 93

TEXT: In order to collect evidence for the hypothesis of surface absorption of low and medium-energy nucleons by nuclei (Bjorklund F. E., Fernbach S. Phys. Rev., 10, 1295, 1958) the authors studied the collision of 6.6-Mev protons with  $F^{19}$  nuclei. Such collisions involve several processes: elastic and inelastic scattering,  $(p,n)$ ,  $(p,\alpha)$ , and  $(p,\gamma)$ . Nine levels with energies between 0.110 and 4.036 Mev are excited by such processes. The total cross section of inelastic scattering of 6.6-Mev protons from  $F^{19}$  nuclei as found from the level excitation cross sections is approximately 450 mb. Data on the cross sections of the other processes were taken from publications. The total absorption cross section is ✓

Card 1/2

Cross section of 6.6-Mev...

S/168/62/000/004/010/010  
B108/B102

$$\sigma_r = \sigma_{p,p} (450 \text{ mb}) + \sigma_{pp} (20 \text{ mb}) + \sigma_{pa} (10 \text{ mb}) + \sigma_{pn} (55 \text{ mb}).$$

As the accuracy of the (p,n) and (p,p) reaction cross sections is not known it is difficult to estimate the error in the total absorption cross section. The exact value of  $\sigma_r$  will certainly be between 500 and 550 mb. ✓

ASSOCIATION: NIYyAF

SUBMITTED: March 14, 1962

Caru 2/2

S/056/62/043/003/001/063  
B125/B102

AUTHORS: Vasil'yev, S. S., Komarov, V. V., Popova, A. M.

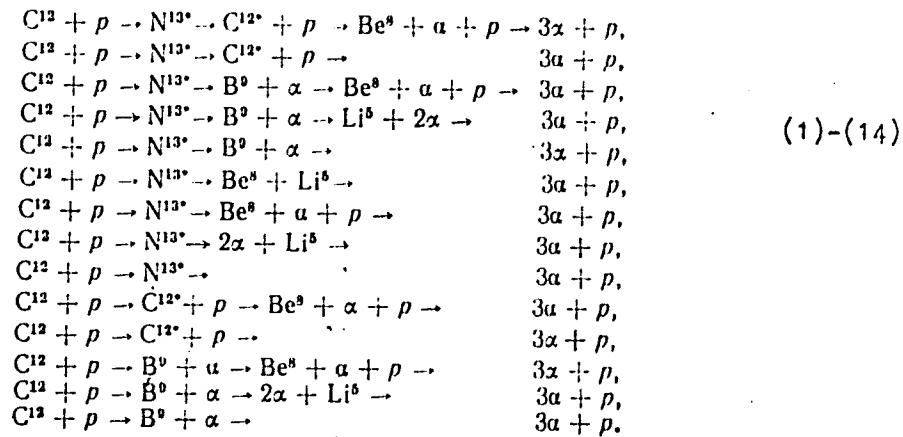
TITLE: Study of decay reactions of carbon and oxygen nuclei under the action of 15-29-Mev protons

PERIODICAL: Zhurnal eksperimental'noy i teoreticheskoy fiziki, v. 43, no. 3(9), 1962, 737-748

TEXT: The energy distribution of the products of the reactions  $C^{12}(p,p'3\alpha)$  and  $O^{16}(p,p'4\alpha)$  was determined from 5 and 6-pronged stars in photographic plates. These plates were irradiated in the 120-cm synchrocyclotron of the Institut yadernoy fiziki MGU (Institute of Nuclear Physics of MGU). For  $C^{12}(p,p'3\alpha)$  reactions and with  $15 \text{ Mev} \leq E_p \leq 29 \text{ Mev}$  the following reaction mechanisms are possible:

Card 1/4

Study of decay reactions of ...

S/056/62/043/003/001/063  
B125/B102

Card 2/4

Study of decay reactions of ...

S/056/62/043/003/001/063  
B125/B102

In these reactions Be<sup>8</sup> may be formed. At excitation energies of the Be<sup>8</sup> nucleus from 0 to 15 Mev only the wide levels  $2.9 \pm 1.5$  Mev and 11.3 Mev occur. At this  $E_p$  of the C<sup>12</sup> nuclei there is only a very low decay probability, according to the mechanisms (1), (2) and (10), (11), with production of the first compound nucleus C<sup>12\*</sup>. In the reaction C<sup>12</sup>(p,p'3 $\alpha$ ) the probability that B<sup>9</sup> will be produced as the first compound nucleus is 20% (mechanisms (12), (13) and (14)). The probability of the mechanisms (3) and (5) is low. The hypothetic mechanisms for the direct decay of the compound system N<sup>13\*</sup> in 2, 3 or 4 noninteracting particles are not confirmed by the experimental energy distributions. At  $15 \text{ Mev} \leq E_p \leq 29 \text{ Mev}$  the reaction C<sup>12</sup>(p,p'3 $\alpha$ ) very probably proceeds by way of a direct nuclear decay of a compound system into three  $\alpha$  particles and one proton. These final particles react in pairs. In the reaction O<sup>16</sup>(p,p'4 $\alpha$ ) the compound nucleus F<sup>17</sup> decays into 4 $\alpha$ +p. These particles resonance-interact in the ground states of the nuclei Li<sup>7</sup> (ground-state) and Be<sup>8</sup> (ground-state, 30-40% probability; excited states  $2.9 \pm 1.5$  Mev, 50%;  $11.3 \pm 0.3$  Mev, 10-20% probability). There are 8 figures and 2 tables.

Card 3/4

Study of decay reactions of ...

S/056/62/043/003/001/063  
B125/B102

ASSOCIATION: Institut yadernoy fiziki Moskovskogo gosudarstvennogo  
universiteta (Institute of Nuclear Physics of the Moscow  
State University)

SUBMITTED: December 20, 1961

Card 4/4

CHESUNOV, V.M., aspirant; VASIL'YEV, S.S., prof., doktor khimicheskikh nauk

Studying the kinetics of ethyl alcohol evaporation from a free surface into a closed gas filled space in connection with the gas passage. Nauch.trudy MTIIP no.18:156-166 '60. (MIRA 15:2)

1. Kafedry teplotekhniki i fiziki Moskovskogo tekhnologicheskogo instituta legkoy promyshlennosti.  
(Ethyl alcohol) (Evaporation)

MENTSOV, V. S., inzh.; VASIL'YEV, S. S., doktor khimicheskikh nauk, prof.

Use of high-frequency current for gluing fabrics. Izv.vys. ucheb.  
zav.; tekhn.leg.prom. no.4:113-118 '61. (MIRA 14:10)

1. Moskovskiy tekhnologicheskiy institut legkoy promyshlennosti.  
Rekomendovana kafedroy fiziki.  
(Gluing)  
(Induction heating)

SHEYNIS, Ye.S., kand.tekhn.nauk, dotsent; VASIL'YEV, S.S., doktor khimi-  
cheskikh nauk, prof.

Effect of high-frequency currents on the reactions of strained  
leather. Nauch.trudy MTILP no.18:52-60 '60. (MIRA 15:2)

1. Kafedra fiziki Moskovskogo tekhnologicheskogo instituta legkoy  
promyshlennosti.

(Leather--Testing) (Strains and stresses)

VASIL'YEV, S.S.; KOMAROV, V.V.; KOSHELYAYEV, G.V.; POPOVA, A.M.

Production of proton beams of different energies within a synchrocyclotron chamber for medium energies. Prib. i tekhn. eksp. 6 no.1:  
17-18 Ja-F '61. (MIRA 14:9)

1. Nauchno-issledovatel'skiy institut yadernoy fiziki Moskovskogo  
gosudarstvennogo universiteta.  
(Protons) (Cyclotron)

VASIL'YEV, S.S.; ROMANOVSKIY, Ye.A.; TIMUSHEV, G.F.

Lower excited states of Ca<sup>40</sup>. Vest. Mosk. un. Ser. 3: Fiz.,  
astron. 16 no. 6:88-89 N-D :61. (MIRA 14:12)

1. Nauchno-issledovatel'skiy institut yadernoy fiziki Moskovskogo  
gosudarstvennogo universiteta.

(Quantum theory)  
(Calcium)

VASIL'YEV, S.S.; ROMANOVSKIY, Ye.A.; TIMUSHEV, G.F.

Collective excited states in  $Mn^{55}$ , Vest. Mosk. un. Ser. 3: Fiz.,  
astron. 16 no.6:89-90 N-D '61. (VFA 14:12)

1. Nauchno-issledovatel'skiy institut yadernoy fiziki Moskovskogo  
gosudarstvennogo universiteta,

(Quantum theory)  
(Manganese)

BASKOVA, K.A.; VASIL'YEV, S.S.; NO SEN CHAN; SHAVTVALOV, L.Ya.

Decay scheme of Br<sup>75</sup>. Zhur. eksp. i teor. fiz. 41 no.5:1484-1486  
N '61.  
(MIRA 14:12)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo  
universiteta.

(Bromine---Decay)

VASIL'YEV, S. S.

Ekonomika Torgovli [by] S.S. Vasil'yev [i Dr] Red.  
koleegiya: S.S. Vasil'yev, G. S. Grigor'yan [i] A.I.  
Fefilov. Moskva, Gostorgizdat, 1962.  
727 p. Tables.

VASIL'YEV, S.S.; ROMANOVSKIY, Ye.A.; TIMUSHEV, G.F.

Inelastic proton scattering on F<sup>19</sup> nuclei. Zhur.eksp.i teor.fiz.  
41 no.4:1040-1042 O '61. (MIFI 14:16)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta.  
(Protons--Scattering)

VASIL'YEV, S.S.; KUMAROV, V.V.; POPOVA, A.M.

Study of the reaction C<sup>12</sup> ( $\alpha$ ,  $4d$ ). Zhur. ekspl. i teor. fiz. 41  
no.6:1757-1760 D '61. (MIRA 15:1)  
(Nuclear reactions) (Carbon)

VASIL'YEV, S.S.; ROMANOVSKIY, Ye.A.; TIMUSHEV, G.F.

Inelastic scattering of 6.6 Mev. protons on Ca<sup>40</sup> and Mn<sup>55</sup> nuclei  
[with summary in English]. Zhur. eksp. i teor. fiz. 42 no.2:395-  
402 F '62. (MIRA 15:2)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta.  
(Protons—Scattering)(Calcium)(Manganese)

BASKOVA, K.A.; VASIL'YEV, S.S.; NO SEN CHAN; SHAVTVALOV, L.Ya.

Study of some radioactive nuclei in the region of the filled  
1 f<sub>7</sub> shell [with summary in English]. Zhur. eksp. i teor. fiz.  
42 no.2:416-426 F '62. (MIRA 15:2)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo  
universiteta.

(Radioisotopes)

VASIL'YEV, S.S.; MIKHALEVA, T.N.; RUDENKO, N.P.; SEVAST'YANOV, A.I.;  
ZAZULIN, V.S.

Long-lived isotope Al26 in structural aluminum used in a nuclear  
reactor. Atom. energ. 11 no.4:401-403 O '61. (MIRA 14:9)  
(Aluminum--Isotopes) (Nuclear reactors)

VASIL'YEV, S.S.; NO SEN CHAN; SHAVTVALOV, L.Ya.

Study of Mn<sup>56</sup> radiation. Izv. AN SSSR. Ser. fiz. 25 no.9:1115-  
1116 '61. (MIRA 14:8)

1. Nauchno-issledovatel'skiy institut yadernoy fiziki Moskovskogo  
gosudarstvennogo universiteta im. M.V. Lomonosova.  
(Manganese--Isotopes)  
(Radiation)

36410  
S/137/62/000/003/105/191  
A060/A101

18. P10C

AUTHORS: Artsishevskiy, M. A., Vasil'yev, S. S., Koshelyayev, G. V.,  
Selisskiy, Ya. P.

TITLE: Action of deuteron irradiation upon the electric resistance of  
alloys undergoing ordering and aging

PERIODICAL: Referativnyy zhurnal, Metallurgiya, no. 3, 1962, 6, abstract 3138  
("Sb. tr. Tsentr. n.-i. in-t chernoy metallurgii", 1959, no. 22,  
168-176)

TEXT: The effect of deuteron irradiation upon the electric resistance R  
of alloys Ni<sub>3</sub>Fe, Fe<sub>3</sub>Al undergoing ordering and of an alloy of Fe with 35% Ni  
and 4.5% Ti undergoing aging was investigated. The specimens were irradiated in  
a cyclotron with deuterons having an energy of 4 Mev. The thickness of the  
specimens constituted 20 - 30 %. The R measurement was carried out by the  
potentiometric method. Because of the small dimensions of the specimens the  
voltage and the current leads constituted a single whole with the working part.  
The specimens of Ni<sub>3</sub>Fe and of Fe<sub>3</sub>Al were investigated in the ordered and the  
unordered states; the specimens of Fe-Ni-Ti - in the aged and hardened states.

Card 1/2

S/137/62/000/003/105/191

A060/A101

Action of deuteron irradiation ...

It was established that when the ordered  $\text{Fe}_3\text{Al}$  alloy is irradiated its R is increased considerably, and the R of the hardened alloy - drops. The bombarding of the  $\text{Ni}_3\text{Fe}$  alloy in the ordered and unordered states causes a considerable decrease in R. In all cases irradiation in fluxes up to  $5 \cdot 10^{17}$  deuterons per  $1 \text{ cm}^2$  causes a sharp change in R, at a further increase of the total flux the rate of change of R drops. The effects uncovered in the Fe-Ni-Ti alloy do not exceed the limits of experimental errors. It is considered that the most probable process causing the reduction in R is the ordering. A considerable drop in the R of the alloy  $\text{Ni}_3\text{Fe}$  is noted, whose degree of ordering corresponds to a temperature of  $250 - 300^\circ\text{C}$ . In this alloy a further occurrence of ordering under irradiation is possible. The shape of the R curves of the irradiated specimens tempered at  $250^\circ\text{C}$  confirms the hypothesis as to the attainment of an intermediate degree of ordering as result of the irradiation. In tempering the  $\text{Ni}_3\text{Fe}$  the soaking time of the specimens at the respective temperatures was insufficient to obtain an equilibrium. The character of the R variation of an irradiated unordered specimen is close to the R variation of an unirradiated ordered specimen. In tempering the Fe-Ni-Ti alloy no great difference in the behavior of irradiated and unirradiated specimens was discovered.

A. Rusakov

[Abstracter's note: Complete translation]

Card 2/2

Coagulating action of alcohols on protein solutions and the Traube rule. B. S. VASIL'YEV.  
 (J. Phys. Chem. Russ., 1929, 13, 43-53).—Coagulation of egg-albumin by the primary alcohol  $C_2-C_6$  was measured. It cannot be due to dehydration as assumed by Kruyt since the coagulating power increases, and the affinity to  $H_2O$  decreases, with rising mol. wt. A real coagulation by dehydration, e.g., with  $(NH_4)_2SO_4$ , is reversible whilst the effect of alcohols is irreversible. The validity of the Traube rule indicates an adsorption mechanism for the coagulation. The amount of the alcohols adsorbed by proteins is calc. J. J. R.

P-1

## Div Organic Chem., Chem Dept. VIEN

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

VASIL'YEV, S. S.

USSR/Chemistry, Colloid - Proteins

Jul/Aug 52

"The Connection Between the Constitution of Organic Compounds and Their Coagulating Effect on Solutions of Egg Albumin," A. P. Sychev, S. S. Vasil'yev, Chair of Phys and Colloid Chem, Moscow Petroleum Institute Acad I. M. Gubkin

"Kolloid Zhur" Vol XIV, No 4, pp 260-266

Dstd the coagulating effect of monocarbinols, polyhydric alcs, phenols, aromatic alcs, alicyclic alcs, aliphatic ketones, alicyclic ketones, aldehydes (including formaldehyde) and aldehyde alcs on solns of dialyzed egg albumin. Found that addn of one

225T13

CH<sub>2</sub>, CH<sub>3</sub>, or Cl group in any series increases the coagulating effect by a factor of 2-2.5; that the relative position of substituents in mono- and polyphenols influences the coagulating effect in a definite manner; etc. Believe that discovery of relationships of this type between the constitution of org compds and their effect on proteins will clarify problems connected with the biol action of org compds.

225T13

VASIL'YEV, S.S.

Chemical Abst.  
Vol. 48 No. 8  
Apr. 25, 1954  
Organic Chemistry

(2) ✓ Relation between the structure of organic compounds and  
their coagulating action on egg albumin solutions. A. P.  
Svchey and S. S. Vasil'ev. *Colloid J. (U.S.S.R.)* 14,  
285-07(1952)(Engl. translation).—See C.A. 47, 2127a.  
H. L. H.

5(

SOV/69-21-2-4/22

AUTHORS: Vasil'yev, S.S. and Yushina, V.V.

TITLE: The Stabilizing Effect of Silver Ions on Albumin (Stabiliziruyushcheye deystviye ionov serebra na al'bumin)

PERIODICAL: Kolloidnyy zhurnal, 1959, Nr 2, pp.148-150 (USSR)

ABSTRACT: This article deals with the effects of silver ions introduced into albumin solutions. After the adsorption of small quantities of silver, the stability of albumin solutions with regard to heating is increased. If larger quantities of silver are adsorbed, the ions produce a coagulating effect. Coagulation takes place, when 30 silver ions per albumin molecule have been adsorbed by the solution. There are 2 graphs and 3 references, 2 of which are German and 1 Soviet.

ASSOCIATION: Tekhnologicheskiy institut legkoy promyshlennosti, Moskva  
(Technological Institute of Light Industry, Moscow)

SUBMITTED: December 12, 1957

Card 1/1

BOCHAGOV, B.A.; VASIL'YEV, S.S.; SEMENCHUK, G.G.; SOLYAKIN, G.Ye.

Energy characteristics of fragments resulting in the fission of Th<sup>232</sup>  
and U<sup>238</sup> nuclei by charged particles. IAd. fiz. 1 no.3:461-470 Mr  
'65. (MIRA 18:5)

1. Fiziko-tehnicheskiy institut im. A.F.Ieffe AN SSSR i Institut  
yadernoy fiziki Moskovskogo gosudarstvennogo universiteta.

BASKOVA, K.A.; VASIL'YEV, S.S.; KHAMO-LEYLA, M.A.; SHAVTVALOV, L.Ya.

Radiations from  $W^{187}$  and  $Tl^{200}W^{187}$  ( $T \approx 24$  hrs).  
Izv. AN SSSR. Ser. fiz. 27 no.10:1258-1260 O '63.

(MIRA 16:10)

VASIL'YEV, S.S.; SHAVTVALOV, L.Yu.

Gamma radiations from  $Au^{197*}$  and the  $\beta^+$ -spectrum of  $O^{15}$ .  
Izv. AN SSSR. Ser. fiz. 27 no.10:1261-1262 O '63.

(MIRA 16:10)

1. Nauchno-issledovatel'skiy institut yadernoy fiziki Moskovskogo  
gosudarstvennogo universiteta im. M.V. Lomonosova.

MEMTSOV, V.S., starshiy prepodavatel'; VOLODINA, N.Ya.; VASIL'YEV, S.S.,  
doktor khim. nauk, prof.

Kinetics of the gluing of shoe materials by means of high-  
frequency currents. Nauch. trudy MTILP no.24:69-73 '62.  
(MIRA 16:7)

1. Kafedra fiziki Moskovskogo tekhnologicheskogo instituta  
legkoy promyshlennosti.

(Gluing) (Induction heating)  
(Shoe manufacture)

GRIGOR'YAN, G.S., prof.; KISTANOV, Ya.A., prof.; FEFILOV, A.I., dots.;  
GENKINA, L.S., dots.; VASIL'YEV, S.S., dots.; SEREBRYAKOV, S.V.,  
prof.; DNEPROVSKIY, S.P., prof.; PIROGOV, P.V., dots.; GOGOL',  
B.I., doktor ekon. nauk; SHOTRINA, N.A., dots.; KULIKOV, A.G.,  
prof.; KUZIN, N.I., dots.[deceased]; AVETISYAN, Ye., red.;  
MUKHIN, Yu., tekhn. red.

[Economics of Soviet trade] Ekonomika sovetskoi torgovli;  
uchebnik. 2., dop. izd. Moskva, Politizdat, 1963. 519 p.

(MIRA 16:12)

(Russia--Commerce)

VASIL'YEV, S.S., dots.; GENKINA, L.S., dots.; GRIGOR'YAN, G.S., dots.;  
KISTANOV, Ya.A., dots.; KULIKOV, A.G., dots.; LIFITS, M.M.,  
prof. [deceased]; OBLOVATSKIY, F.Ya., dots.; PIROGOV, P.V., dots.;  
POPOV, A.N., dots.; SNOTRINA, N.A., dots.; FEFILOV, A.I.,  
STARCKHOVA, I.I., red.; EL'KINA, E.M., tekhn. red.

[Economics of commerce] Ekonomika torgovli. Ned. kollegija;  
Vasil'ev, S.S., Grigor'yan, G.S., Fefilov, A.I. Moskva, Gos-  
torgizdat, 1962. 727 p. (MIRA 15/6)  
(Commerce)

VASIL'YEV, S.S.; ROMANOVSKIY, Ye.A.; TIMUSHEV, G.F.

Measuring the angular distribution for the  $\text{Al}^{27} (\text{p}, \text{p}')\text{Al}^{27*}$  reaction with the aid of a magnetic analyzer when  $E_p = 6.6$  Mev.  
*Zhur.eksp.i teor.fiz.* 40 no.3:972-973 Mr '61. (MIRA 14:8)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta.

(Nuclear reactions) (Magnetic measurements)  
 (Aluminum—Isotopes)

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

S/048/61/025/009/002/007  
 B104/B102

246400

AUTHORS: Vasil'yev, S. S., Komarov, V. V., and Popova, A. M.

TITLE: Properties of the lower states of the  $\text{Li}^5$  and  $\text{Be}^8$  nuclei produced in decays of light nuclei

PERIODICAL: Akademiya nauk SSSR. *Izvestiya. Seriya fizicheskaya*, v. 25, no. 9, 1961, 1117 - 1120

TEXT: This paper was read at the 9th Annual Conference on Nuclear Spectroscopy. It deals with the question how the influence of interaction of reaction products in the final state can be taken into account when studying the integral energy distribution of particles produced by direct decay of a compound nucleus. The differential cross section of the decay of a compound nucleus with spin I and parity  $\pi$  into n particles with the energies  $E_i$  and the momenta  $\vec{p}_i$  can be written as

$$d\sigma \sim \delta \left\{ \sum_{i=1}^n (E_i - E) \right\} \delta \left( \sum_{i=1}^n p_i \right) \prod_{i=1}^n d^3 p_i |H_{ab}|^2 ,$$

27477

S/048/61/025/009/002/007

B104/B102

Properties of the lower states of ...

where  $E$  is the total energy of the system, and  $H_{ab}$  is the matrix element of the transition from the initial state  $a$  into the final state  $b$ . This element has the form

$\langle \prod_{i=1}^n \psi_i \chi(s_i) | v | \Psi_{jjz} \rangle$ ,  $\Psi_{jjz}$  being the wave function of the compound nucleus and  $\psi_i \chi(s_i)$  the wave and spin function of the  $i$ -th particle. If the particles do not interact in the final state, the  $\psi_i$  represent plane waves. In this case, the motion of two particles,  $i$  and  $j$ , can be represented as the product of two plane waves:  $\exp(-i\vec{q}\cdot\vec{\varphi}/\hbar) \exp(-i\vec{p}\cdot\vec{R}/\hbar)$ , where  $\vec{q} = \frac{1}{2}(\vec{p}_i - \vec{p}_j)$  is the momentum of the relative particle motion. The wave function of the relative motion of the interacting particles  $i$  and  $j$  is a distorted wave. The radial part of the wave function of the relative motion of the scattered particles is approximately given by  $\varphi(\rho) = f(E_{rel})g(\rho)$ , as was shown by E. W. Hamburger (Thesis, University of Pittsburgh, 1959). Using this expression, the differential reaction

Card 2/6

27477

S/048/61/025/009/002/007

B104/B102

Properties of the lower states of ...

cross section is written as

$$\frac{d\sigma}{dE_i} \sim f(E_i) \frac{\sin^2(\delta + \Phi)}{q^2}$$
. If a simultaneous interaction between two pairs of particles in the final state is possible, the differential cross section acquires the form

$$\frac{d\sigma}{dE_i} \sim f(E_i) \frac{\sin^2(\delta_1 + \Phi_1)}{q_1^2} \cdot \frac{\sin^2(\delta_2 + \Phi_2)}{q_2^2} \quad (6).$$

This is illustrated by a study of the energy distribution of protons from the decay of  $C^{12}$  into three alphas induced by 15 - 30 Mev protons. A previous paper by the authors (S. S. Vasil'yev et al., Izv. AN SSSR, Ser. fiz., 24, 1145 (1960)) has shown that four- and three-particle decays of  $C^{12}$  may be accompanied by interactions of two alphas at the levels 0, 2.9, and 11.8 Mev of the  $Be^8$  nucleus, and by an  $\alpha$ - p interaction at the ground level of the  $Li^5$  nucleus. This indicates that  $Li^5$  and  $Be^8$  occur, not as

Card 3/6

Properties of the lower states of ...

27477  
S/048/61/025/009/C02/007  
B104/B102

compound nuclei of a cascade, but as a result of interaction of the decay products. The excitation energies of the Be<sup>8</sup> and Li<sup>5</sup> nuclei should then appear in the histograms, and the alpha and proton spectra should be described by statistical distributions calculated on the assumption of a four-particle decay with the above-mentioned interactions. To prove this assumption, the authors investigated the energy distribution of protons from the C<sup>12</sup>(p,p')3 $\alpha$  reaction at energies of 15 - 30 Mev of the primary protons. (6) is represented in a form corresponding to this case. The resulting curve for the proton energy distribution agrees well with the experimental results. There are 1 figure and 7 references: 5 Soviet and 2 non-Soviet. The references to English-language publications read as follows: Hamburger E. W., Thesis, University of Pittsburgh, 1959; Brueckner K., Phys. Rev., 82, 598 (1951).

ASSOCIATION: Institut yadernoy fiziki Moskovskogo gos. universiteta im. M. V. Lomonosova (Institute of Nuclear Physics of Moscow State University imeni M. V. Lomonosov)

Card 4/6

VASIL'YEV, S.S. (Moskva)

Calculation of the concentration of excited molecules in the zone of stationary electrical discharges. Zhur. fiz. khim. 35 no. 4:761-771 Ap '61.  
(MIRA 14:5)

1. Moskovskiy tekhnologicheskiy institut legkoy promyshlennosti.  
(Molecular dynamics) (Electric discharges through gases)

VASIL'YEV, S.S.; SHAVTVALOV, L.Ya.

$\beta^+$  -spectrum of Si<sup>27</sup>. Zhur.eksp.i teor.fiz. 39 no.5:1221-1223  
N '60. (MIRA 14:4)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo  
universiteta.  
(Silicon--Spectra)

S/120/61/000/001/003/062  
E032/E11<sup>4</sup>

AUTHORS: Vasil'yev, S.S., Komarov, V.V., Koshelyayev, G.V.,  
and Popova, A.M.

TITLE: Production of Proton Beams of Various Energies Inside  
the Synchrocyclotron Chamber at Intermediate Energies

PERIODICAL: Pribory i tekhnika eksperimenta, 1961, No.1, pp.17-18

TEXT: In nuclear reaction studies employing protons of intermediate energies inside the synchrocyclotron chamber, it is convenient to use a method in which a number of targets are simultaneously irradiated by proton beams of approximately equal intensity but different energy (with sufficiently small energy spread in each beam). For this purpose the main proton beam is directed on to an internal target in the form of a wedge. In the latter the original protons are slowed down and scattered in different ways so that the protons leaving the wedge have an energy spectrum. In the magnetic field protons of different energies move over trajectories of different radii. These trajectories are intercepted by a set of slits which thus define a number of proton beams of different energies. The slits are located on the bottom

Card 1/ 4

S/120/61/000/001/003/062  
E032/E114

Production of Proton Beams of Various Energies Inside the Synchrocyclotron Chamber at Intermediate Energies

of the chamber and are arranged in such a way that they let through only those protons which are scattered at small angles in the downward direction but are practically unscattered in the horizontal plane. This method has been used in nuclear reaction studies using the 120 cm synchrocyclotron of the Scientific Research Institute of Nuclear Physics of the Moscow State University (Nauchno-issledovatel'skiy institut yadernoy fiziki MGU) (initial proton energy 30 MeV). The wedge was made of copper and had an angle of 40°. The intercepting slits were 3 mm wide each and defined 9 proton beams in the energy range 7.5-30 MeV. The energy spread in each channel was smaller for the smaller energies. The nine beams were allowed to strike nuclear emulsions at an angle of 6°. In order to obtain approximately equal intensities in the 9 channels the working part of the wedge was made approximately equal to the radial half-width of the synchrocyclotron beam.

There is 1 figure.

Card 2/ 4

S/120/61/000/001/003/062  
E032/E114

Production of Proton Beams of Various Energies Inside the  
Synchrocyclotron Chamber at Intermediate Energies

Fig.

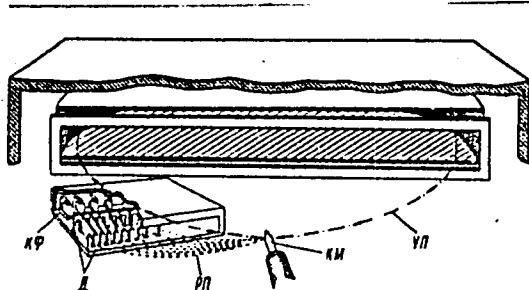


Figure caption: KM - wedge target; YII - main beam;  
YI - scattered protons; A - slits;  
KΦ - emulsions

Card 3/4

S/120/61/000/001/003/062  
E032/E11<sup>4</sup>

Production of Proton Beams of Various Energies Inside the  
Synchrocyclotron Chamber at Intermediate Energies

ASSOCIATION: Nauchno-issledovatel'skiy institut yadernoy fiziki  
MGU  
(Scientific Research Institute of Nuclear Physics,  
MGU)

SUBMITTED: December 10, 1959

Card 4/4

VASIL'YEV, S.V.

Action of nitrogen tetroxide on crotonic acid. Zhur. ob. khim. 30  
no.10:3412-3414 O '61. (MIRA 14:4)

1. Moskovskiy institut tonkoy khimicheskoy tekhnologii  
(Crotonic acid) (Nitrogen oxide)

VASIL'YEV, S.V.; ZHURAVLEVA, A.A.; KOSTOMAROVA V.L.; VASIL'YEV, G.S.

Action of nitrogen tetroxide on dibenzalacetone. Zhur.ob.khim. 30  
no.10:3414-3416 O '61. (MIRA 14:4)

I, Moskovskiy institut tonkoy khimicheskoy tekhnologii.  
(Nitrogen oxide) (Pentadienone)

KHORIN, Vasiliy Ivanovich, kand.ekonom.nauk; VASIL'IEV, S.S., red.;  
PAVLOVA, A.S., red.izd-va

[The turnover of consumers' cooperatives in the postwar period]  
Tovarоoborot potrebitel'skoi kooperatsii v poslevoennyyi period.  
Moskva, Izd-vo TSentrosciiza, 1959. 134 p. (MIRA 13:8)  
(Cooperative societies)

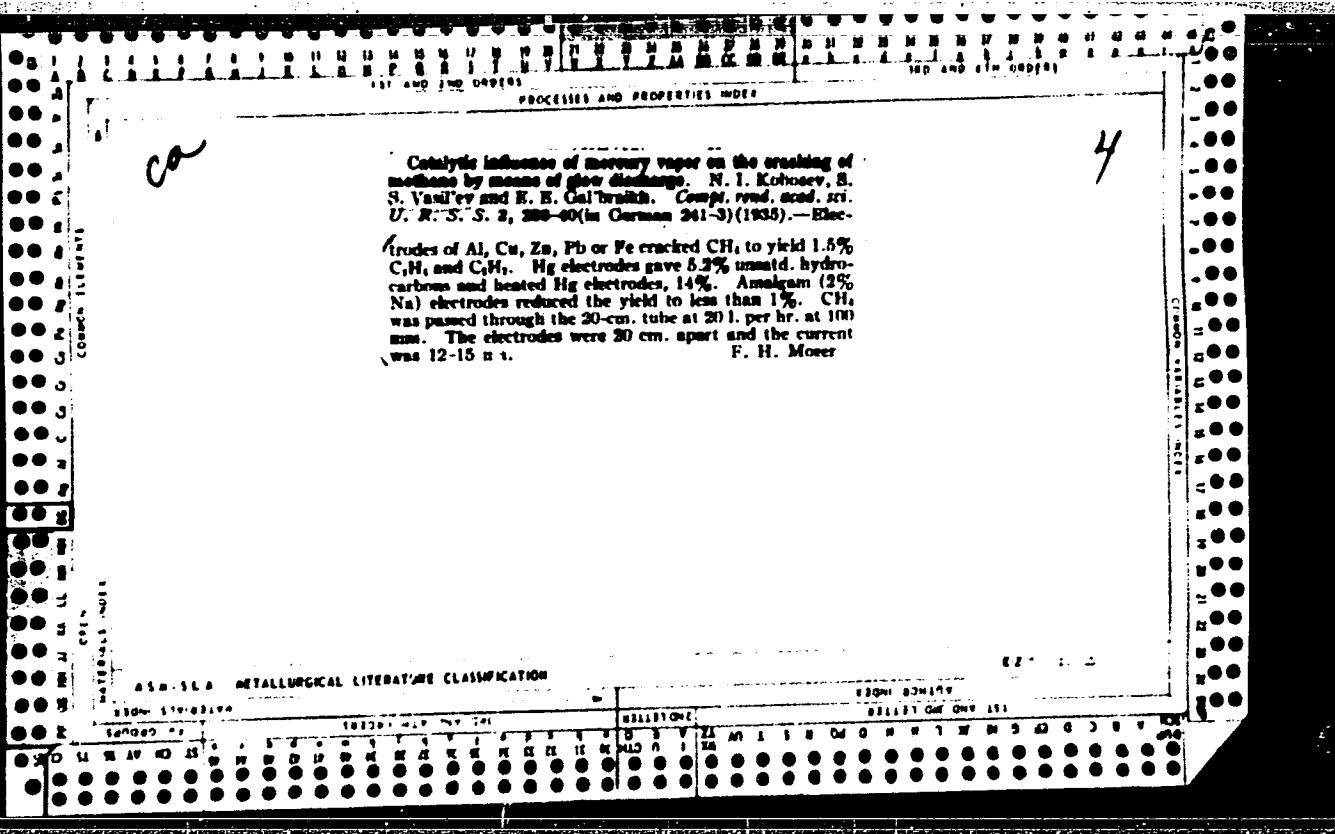
"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

Inversion of heats of melting of powders by various methods. II V. Hlin and  
S. S. Vasil'ev. *J. Phys. Chem. (U. S. S. R.)* 2, 2027 9(1931). - Data on heating-heated  
for various times show a reverse order of heat effect toward various  $\text{MgO}-\text{H}_2\text{O}$  mixts.  
V. H. Rathmann

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"



PROCESS AND PROPERTIES INDEX

*✓ The kinetics of the oxidation of sulfur dioxide solutions.*  
 S. B. Vasilev, L. I. Kashtanov and T. L. Kastorskaya.  
*Acta Physicochim. U. R. S. S.*, 3, 413-414 (1935) (in German);  
*J. Phys. Chem. (U. S. S. R.)* 7 (in press) (1936).—When  
 air bubbles through soln. of  $\text{SO}_2$  in  $\text{H}_2\text{O}$ , the  $\text{SO}_2$  is re-  
 moved from soln. at a rate given by the equation  $\frac{dc}{dt} = r_1 - r_2c$ , where  $c$  is the concn. of  $\text{SO}_2$  in soln.,  $t$  the concn.  
 in the gas-phase bubbles,  $l$  the height of the soln. through  
 which the bubbles pass, and  $r_1 = 0.1$  and  $r_2 = 0.0018$ ,  
 are absorption and desorption constns., resp. The rate of  
 oxidation of  $\text{SO}_2$  solns. by an air stream at  $15^\circ$  is given by  

$$\Delta = \delta/c^x$$
, where  $\delta = \text{mols. oxidized}$ ,  $x$  is an exponent  
 slightly less than unity and decreasing with increasing  $c$ ,  
 and  $\Delta$  is a const. = 7.7. While  $x$  changes from 0.5 to  
 $5 \times 10^{-3} M$  the  $x$  decreases from 0.9 to 0.75 owing to  
 insufficiency of O. On addn. of 0.01%  $\text{MnSO}_4$  to the  $\text{SO}_2$   
 soln. the value of  $\Delta$  for a molar soln. is increased to 35.0  
 but  $x$  takes on much smaller values, decreasing from 0.6  
 to 0.2 over the same range. Math. analysis of the data  
 indicates that the catalytic effect of  $\text{MnSO}_4$  is due to the  
 formation of an unstable intermediate compd. The  
 intensity of oxidation is characterized by the function  

$$r = r_1/(r_1 + a)$$
, where  $a$  is a rate const. for the oxidation  
 reaction. F. H. Rathmann

## ASM-SEA METALLURGICAL LITERATURE CLASSIFICATION

E3001 834177

E3001 834177

E3001 834177 ONE ONLY 151

*CN*

A study of chain processes in the reaction between sulfur dioxide and ozone. I. Stoichiometry in the oxidation of sulfur dioxide with ozone in aqueous solution. S. S. Vasilev, M. V. Frolov, L. I. Kashtanov and T. I. Kostomukha. J. Gen. Chem. (U. S. S. R.) 5, 110-51 (1935). Two series of experiments were made on the oxidation of SO<sub>2</sub> dissolved in H<sub>2</sub>O with dissolved air. In one series, the concn. of O<sub>3</sub> in the air was const. (1.2%), while the concn. of SO<sub>2</sub> in H<sub>2</sub>O varied (0.02, 0.1 and 0.5 N); in the other series, the concn. of SO<sub>2</sub> was const. (0.02 or 0.1 N), while the concn. of O<sub>3</sub> varied (from 0.018 to 1.2%). The "stoichiometric" coeff.  $\alpha$ , defined as the number of mole of oxidized SO<sub>2</sub> per mole of O<sub>3</sub> used up, varied from

1 to 5, depending on the concn. of SO<sub>2</sub> (0.02-0.5 N) at an O<sub>3</sub> concn. of 1%. For each concn. of SO<sub>2</sub> there is a definite concn. of O<sub>3</sub> for which  $\alpha$  is a max. This may vary between 0 and 15, i.e., it was 1 & 2 times as high as found by other workers. The high value of  $\alpha$  leads to the assumption of a chain character of the reaction involved.

S. L. Madorsky

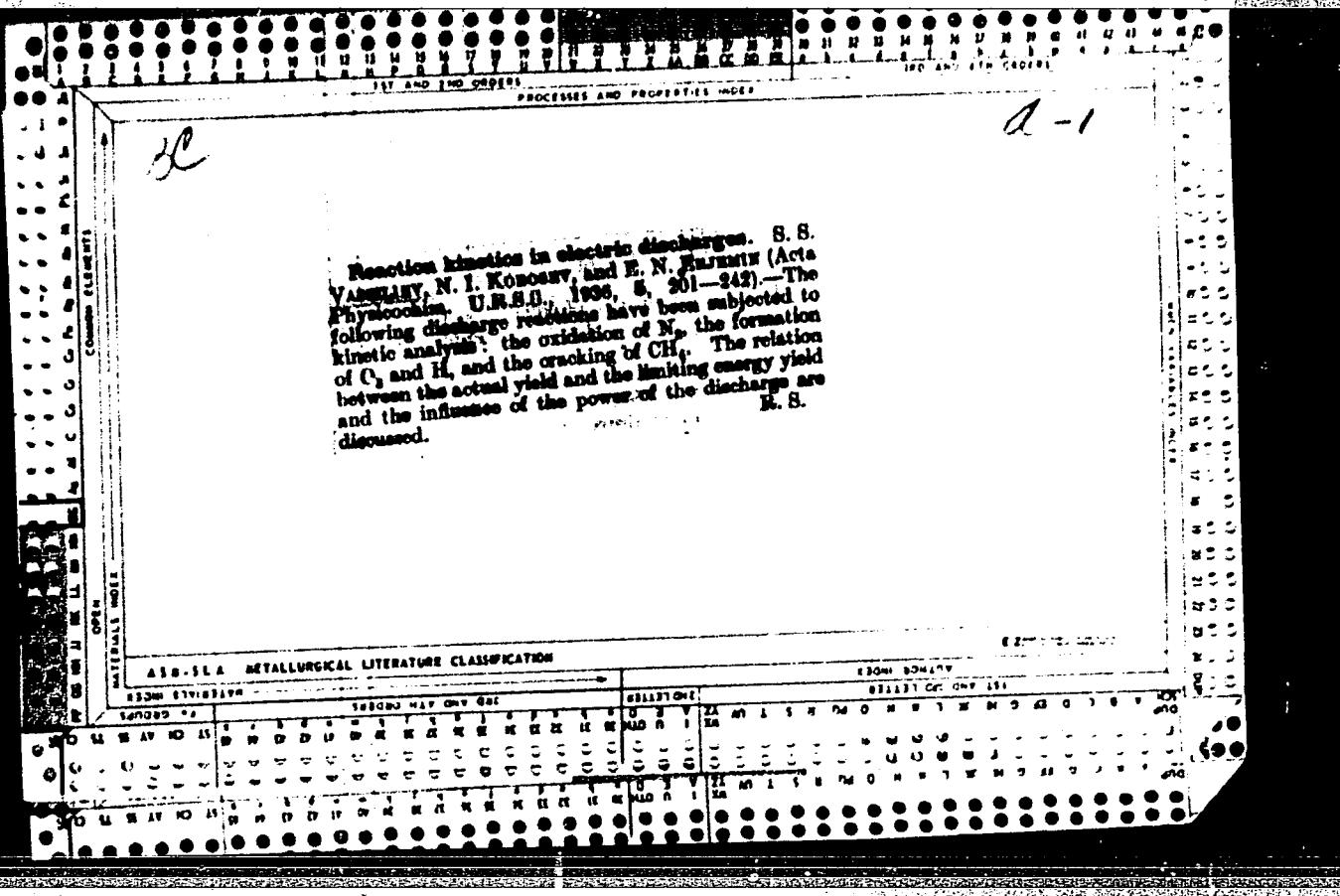
## AIA-314 METALLURGICAL LITERATURE CLASSIFICATION

Thermodynamics of endothermic processes of nitrogen fixation. N. I. KOROBIN, S. S. VASIL'YEV, and J. S. KAAKOVUORI (Acta Physicochim. U.R.S.S., 1934, 4, 245-263).—Equilibrium and energy relations have been calculated thermodynamically at 2000°, 3000°, and 3600° for the reactions  $0.5N_2 + 0.5O_2 = NO$  (a),  $0.5N_2 + CO = NO + CO$  (b),  $0.5N_2 + 0.5H_2 + C = HCN$  (c),  $0.5N_2 + 0.5C_2H_2 = HCN$  (d), and  $0.5N_2 + OH = HCN + 1/2H_2O$  (e). The neglect of resonance and changes such as dissociation gives rise to serious errors, and the results obtained by Kraus and Mackay (A.J. 1933, 1934) for (b) are considered to be incorrect. The reactions (c) and (d) give considerable yields of HCN and require little energy. The considerable yield of NO obtained in the glow discharge is connected with the low temp.

H. H. U.

**APPROVED FOR RELEASE: 08/31/2001**

CIA-RDP86-00513R001858910020-7"



CA

7

**Kinetics of reactions in electric discharges.** S. S. Vassil'ev, N. I. Koborev and F. N. Kremin. *J. Russ. Chem. (U. S. S. R.)* 7, 619-644 (1934). The authors, applying the laws of chem. kinetics, discuss the theory of reactions, reaction velocities, course and yields as a function of discharge intensity and apply their ideas to data from the literature on the oxidation of NO, formation of ozone and of at. H and to the cracking of methane. It is shown that the energy distribution is different for thermal and discharge reactions, that the kinetic constants and the course of the reaction are functions of the power of the discharge.

## 430.14 METALLURGICAL LITERATURE CLASSIFICATION

APPROVED FOR RELEASE: 08/31/2001

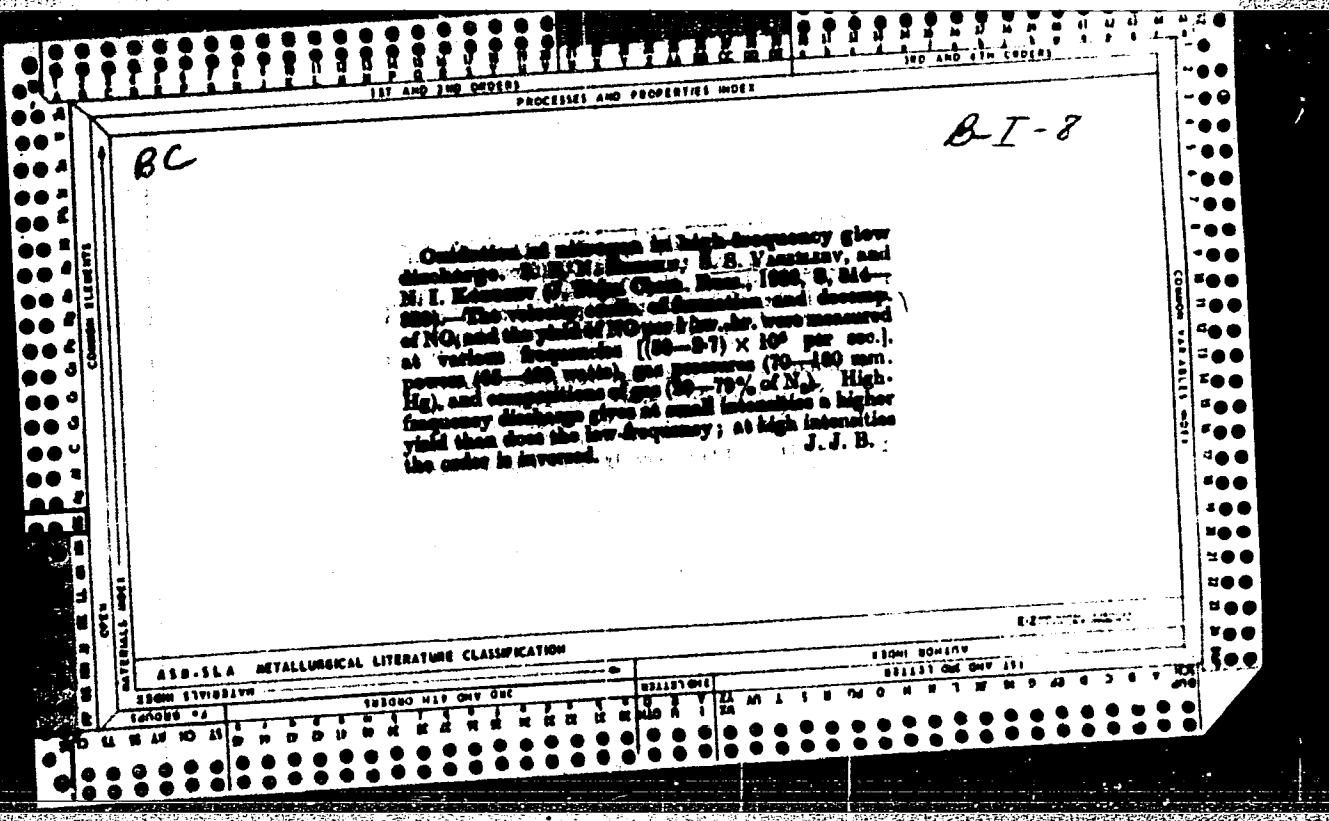
CIA-RDP86-00513R001858910020-7"

## PROCESSES AND PROPERTIES

The equation of state of a gas. S. S. Vasil'ev. *J. Phys. Chem. (U. S. S. R.)* 7, 726-32 (1933). Using the data of Ramsay and Young (*Phil. Trans. A178*, 85 (1887)) for Rn<sub>2</sub> vapors from 320° to 490°K., V. finds that the function  $E = (RT - \rho V)$  where  $R = 841.080$  cm.<sup>3</sup> atm./degree is not a function of  $V$  and is not a linear function of  $T$  but is given by  $E = 10^{10.19}/T^{1.01}$  to within the exp'l. error. The data of Battelle also lead to this equation.

V. H. Rathmann

## APPENDIX METALLURGICAL LITERATURE CLASSIFICATION



CA

21

Application of catalysts and ozone in the water purification of flue gases from sulfur dioxide. S. S. Vasil'ev, L. I. Kashtanov, T. L. Kastorskaya and O. G. Nemtsova. *Vestn. Tekhniki* 1936, No. 58, 9, 12-13. A soln. of Mn salts (0.1%) used for the absorption of SO<sub>2</sub> results in a 25% H<sub>2</sub>SO<sub>4</sub> soln. This may be explained by the formation (under certain conditions) of an active form of catalyst. The action of O<sub>3</sub> in the process of SO<sub>2</sub> oxidation depends on the ratio of its concn. to that of SO<sub>2</sub>. At equal concn. one mol. of H<sub>2</sub>SO<sub>4</sub> is formed per mol. of O<sub>3</sub>, and if there is a high excess of SO<sub>2</sub> then 15-20 mol. of H<sub>2</sub>SO<sub>4</sub> is formed per mol. of O<sub>3</sub>. This is explained by the chain reaction between O<sub>3</sub> and SO<sub>2</sub>. The formation of an active form of the catalyst is probably promoted by O<sub>3</sub>.

A. A. Pudgorny

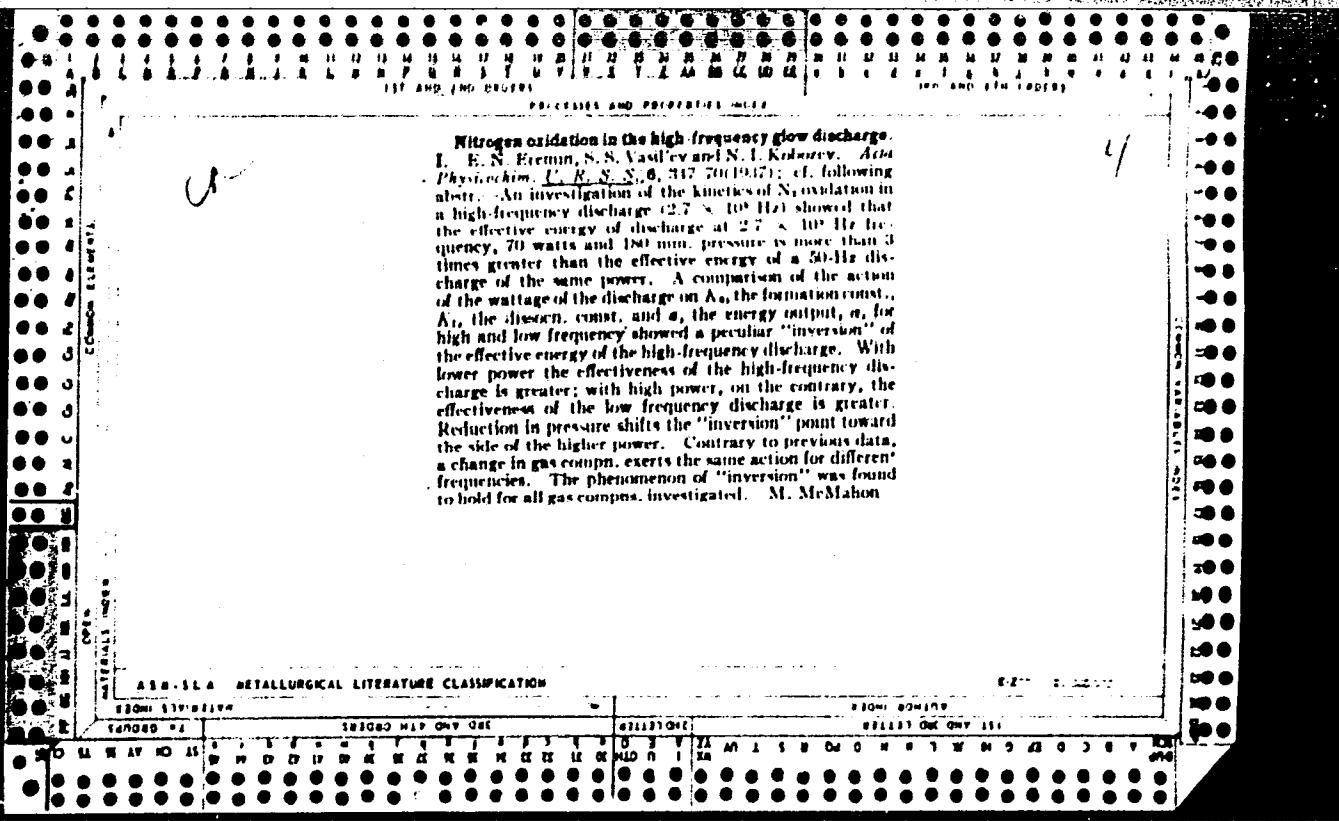
## AIRO-SLA METALLURGICAL LITERATURE CLASSIFICATION

SEARCHED		INDEXED		SERIALIZED		FILED	
■	■	■	■	■	■	■	■
■	■	■	■	■	■	■	■

Preventing the formation of aluminum carbide in the electrolytic production of aluminum. S. Vasil'ev. *Nauk. i Tekhniki* 6, No. 23, 3 (1937); *Chem. Zentr.* 1938, II, 3737.—The formation of Al carbide is promoted by an excess of NaF in the electrolyte and very definitely suppressed when AlF<sub>3</sub> predominates. Carbide formation is accelerated by elevated temps. The carbide formed in the cell can be decompd. by blowing air or O through it.

M. G. Moore

4



"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

Influence of the current density on the yield of acetylene  
in electrical cracking of methane S. S. Vasil'ev, N. I.  
Kolosov and E. N. Efremov. Khim. Treridza. Teplofiz.  
N, 73 No 10(1977). A criticism of the above paper.  
A. A. Podgorny.

ASH-VLA METALLURGICAL LITERATURE CLASSIFICATION

CLASS NUMBER

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

*ca*

Nitrogen oxidation in a high-frequency glow discharge.  
H. E. N. Premin, B. S. Vasil'ev and N. I. Kulozhev.  
*J. Phys. Chem. (U. S. S. R.)* 6, 48-61; *Acta Physicochim.*  
*U. R. S. S.*, 6, 631-64 (1937); cf. preceding abstr.  
—At from 70 to 180 mm. pressure and 70 w., a  
high-frequency ( $2.7 \times 10^9$  Hz) discharge is 3 times as  
efficient as a 50-Hz discharge. Above 90 w., the  
low-frequency discharge is the more effective. A com-  
parison of the effects of  $2.7 \times$ ,  $5 \times$ ,  $7 \times$  and  $10 \times 10^9$  Hz  
frequencies shows that the effect is the same in all cases  
and hence not due to chem. resonance effects. No differ-  
ence in effect was observed when either internal or  
external electrodes were used. At higher wattages the  
max. yield is obtained at higher relative concns. of O<sub>2</sub>.  
Electron temp. as detd. from the spectra of the dis-  
charge is higher at 70 w. high frequency than at low  
frequency and is higher at low wattage than at high  
wattage, and higher at low pressure than at high pressure.  
F. H. Rathmann

ECONOMIC SECURITY ELEMENTS

OPEN

MATERIALS HOLD

## ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

X3001 319-02194

100292 47

VOLUME 417 ONLY Det

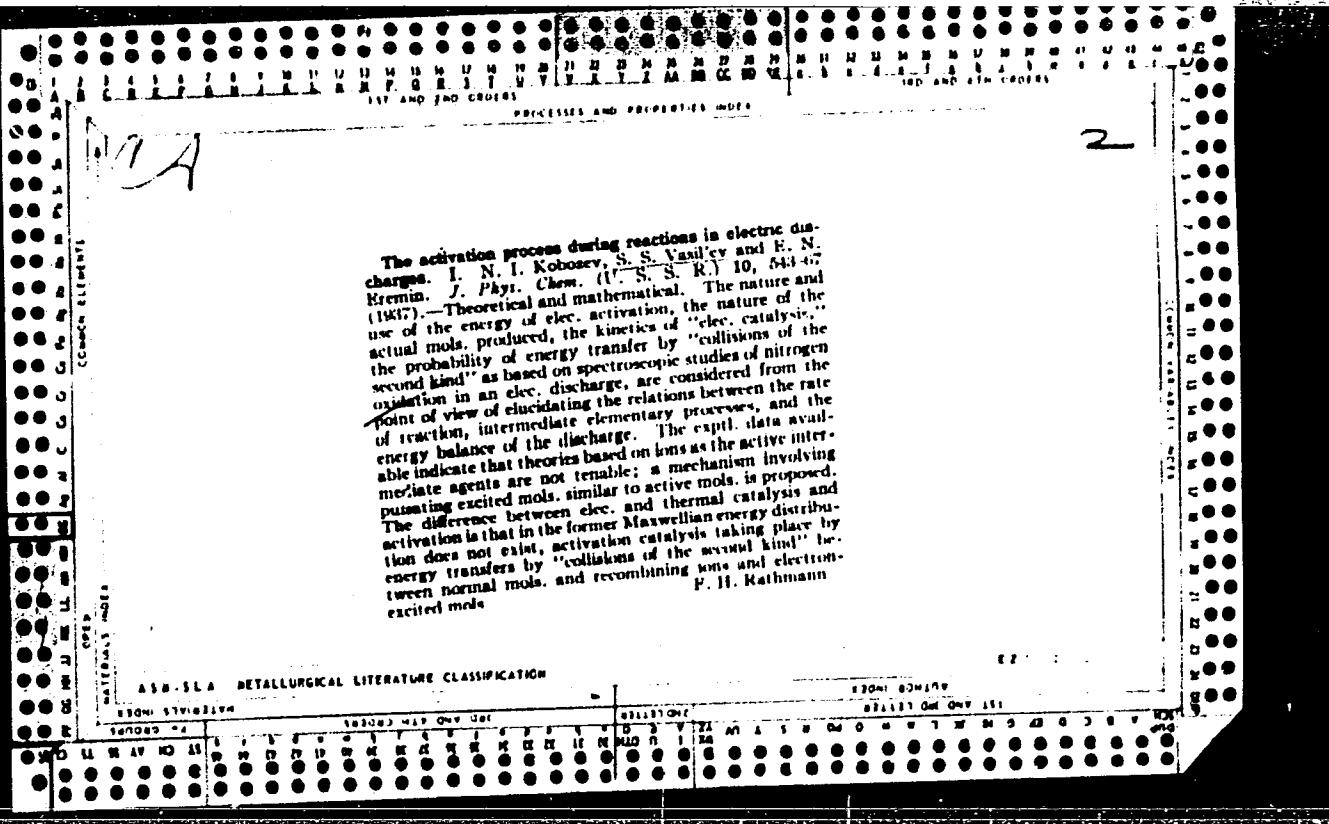
SECTION 1

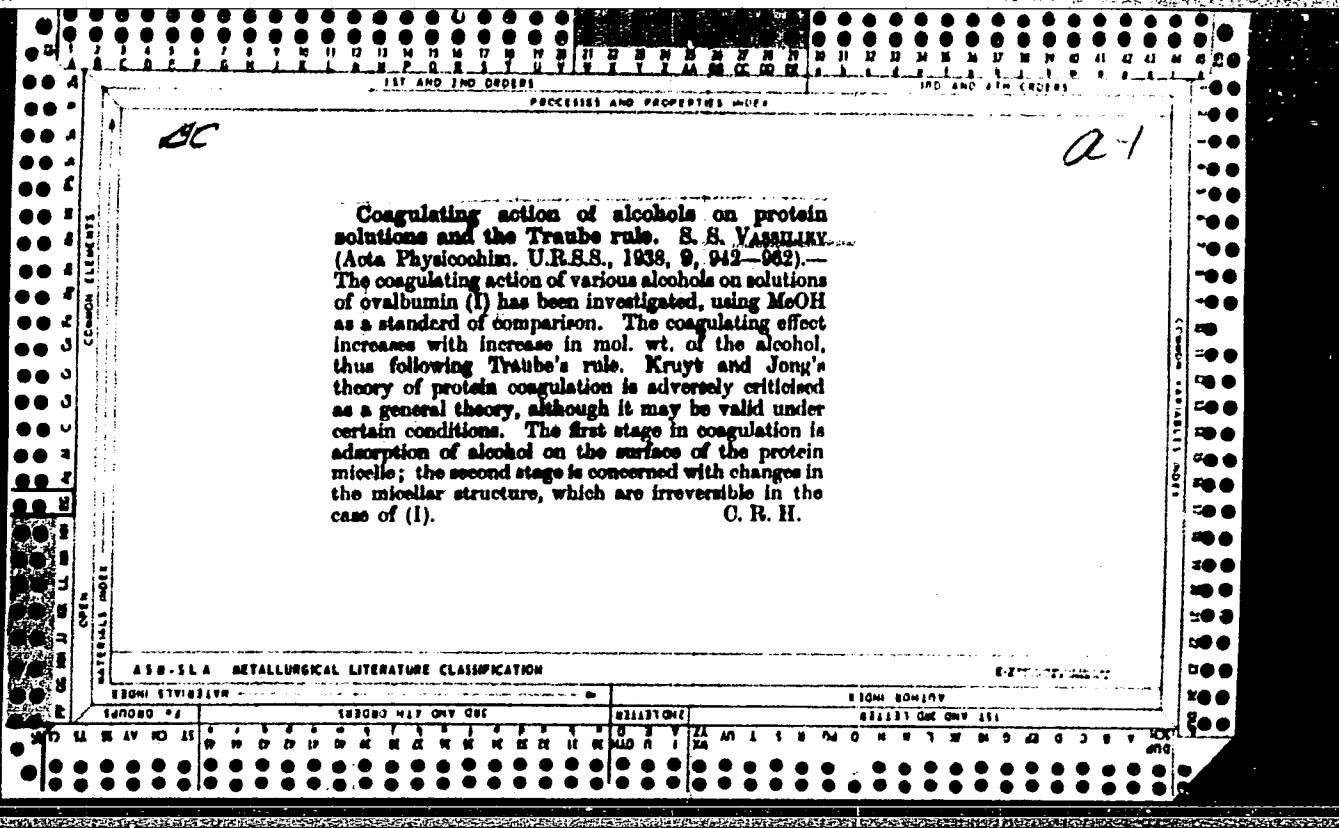
SERIALS 1

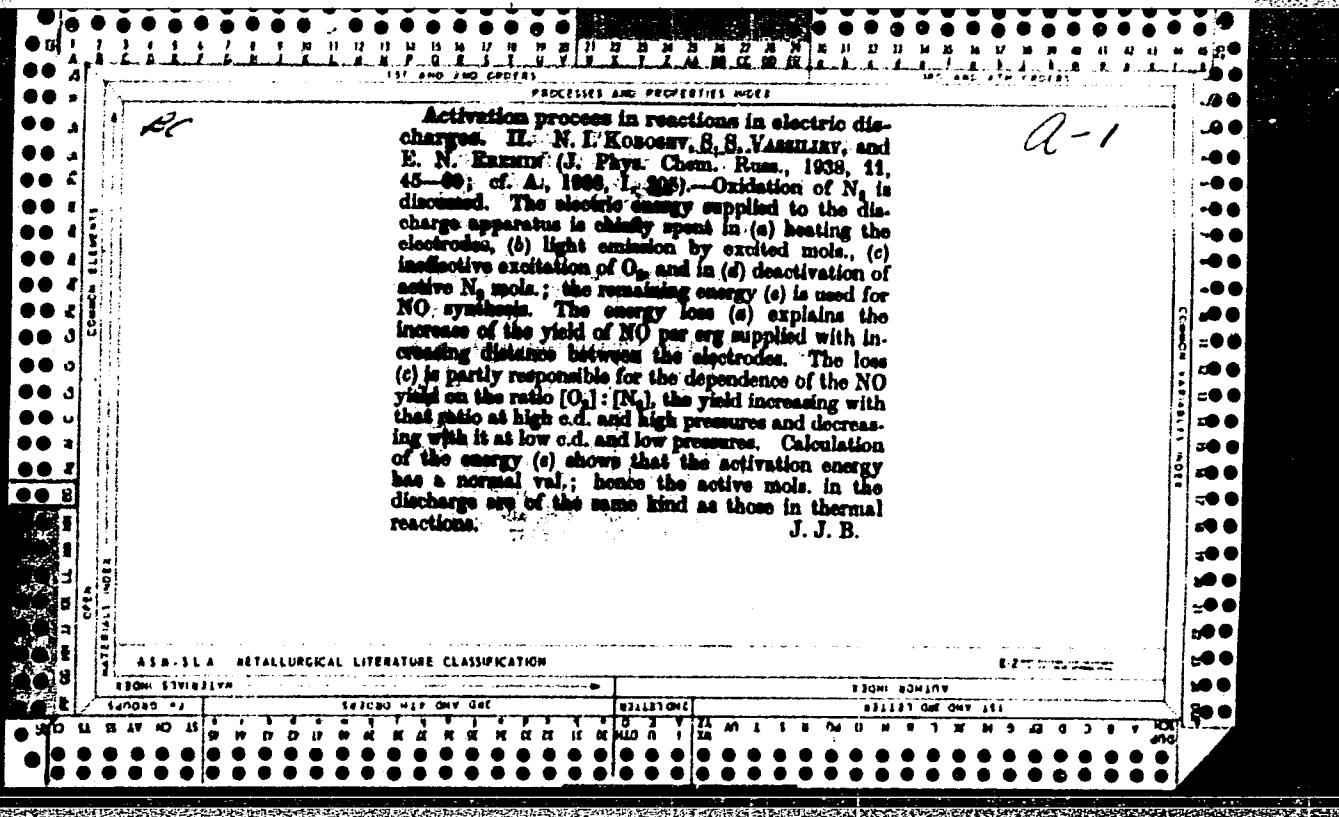
THE QDV 151

The kinetic mechanism of the chain oxidation of sulfur dioxide in solution by oxidized air. S. S. Vasilev, L. I. Kashtanov and T. I. Kuznetsova. *J. Phys. Chem.* (U. S. S. R.) 10, 320 (1937). The chain lengths for the reaction  $\text{SO}_2 + \text{H}_2\text{O} + \text{O}_3 \rightarrow \text{HSO}_3 + \text{O}_2$  involving the stages  $\text{SO}_3$ ,  $\text{HSO}_3$  and  $\text{HSO}_4$  vary from 1.05 for 1.32%  $\text{O}_3$  and 3.0%  $\text{SO}_2$  to 1.7 for 0.03% and 2.0% and 1.0 for 0.04% and 2.0%. In the equation  $W = e^{-1/\theta}$ , where  $\theta$  = chain length,  $e$  = ratio  $\text{SO}_2$  to  $\text{O}_3$ ,  $W = 0.30$  for the reaction as given above, catalyzed by Mn salt (0.01%). The reaction is unimol. with respect to ozone. In strong  $\text{HSO}_4$  the reaction is accelerated 3-4 times. Ozone in low concn. can be successfully used to convert  $\text{SO}_2$  to  $\text{HSO}_4$ .

F. H. Rathmann

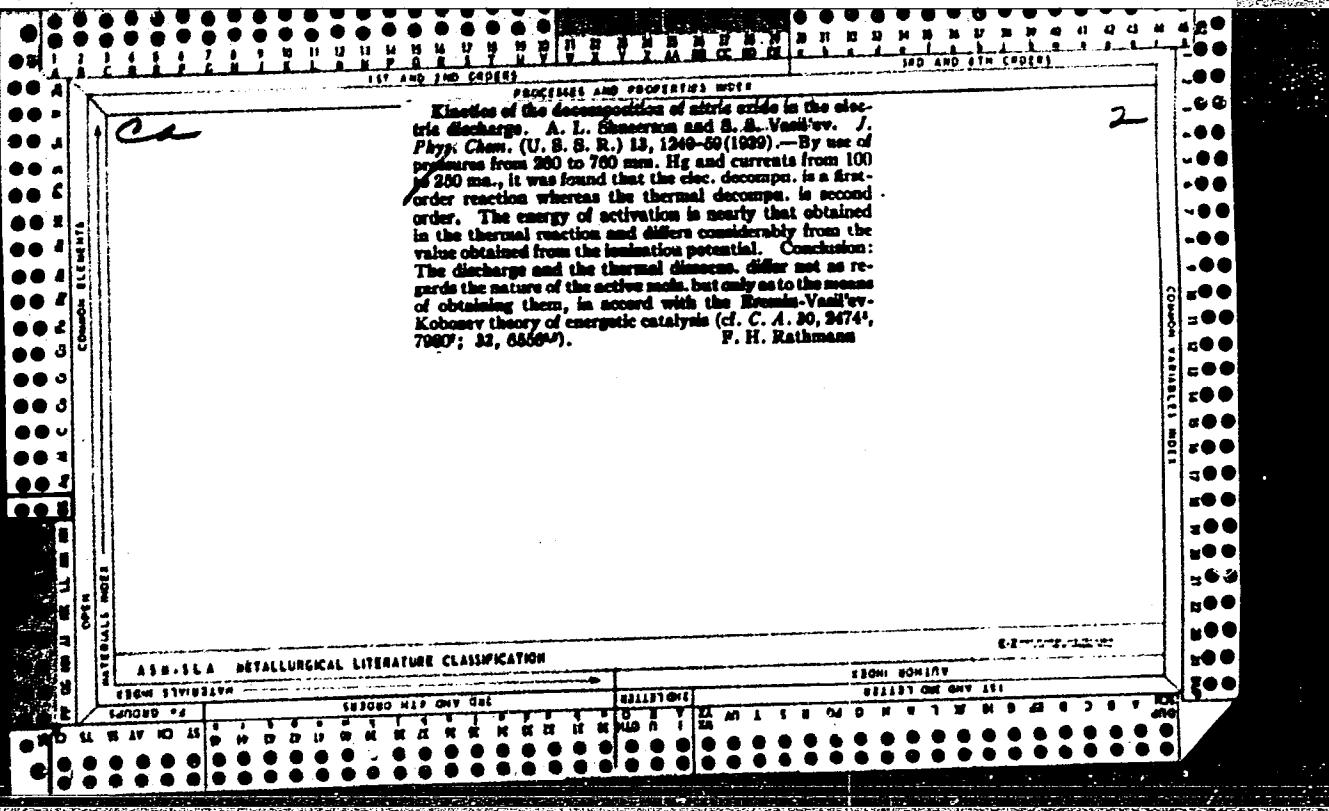


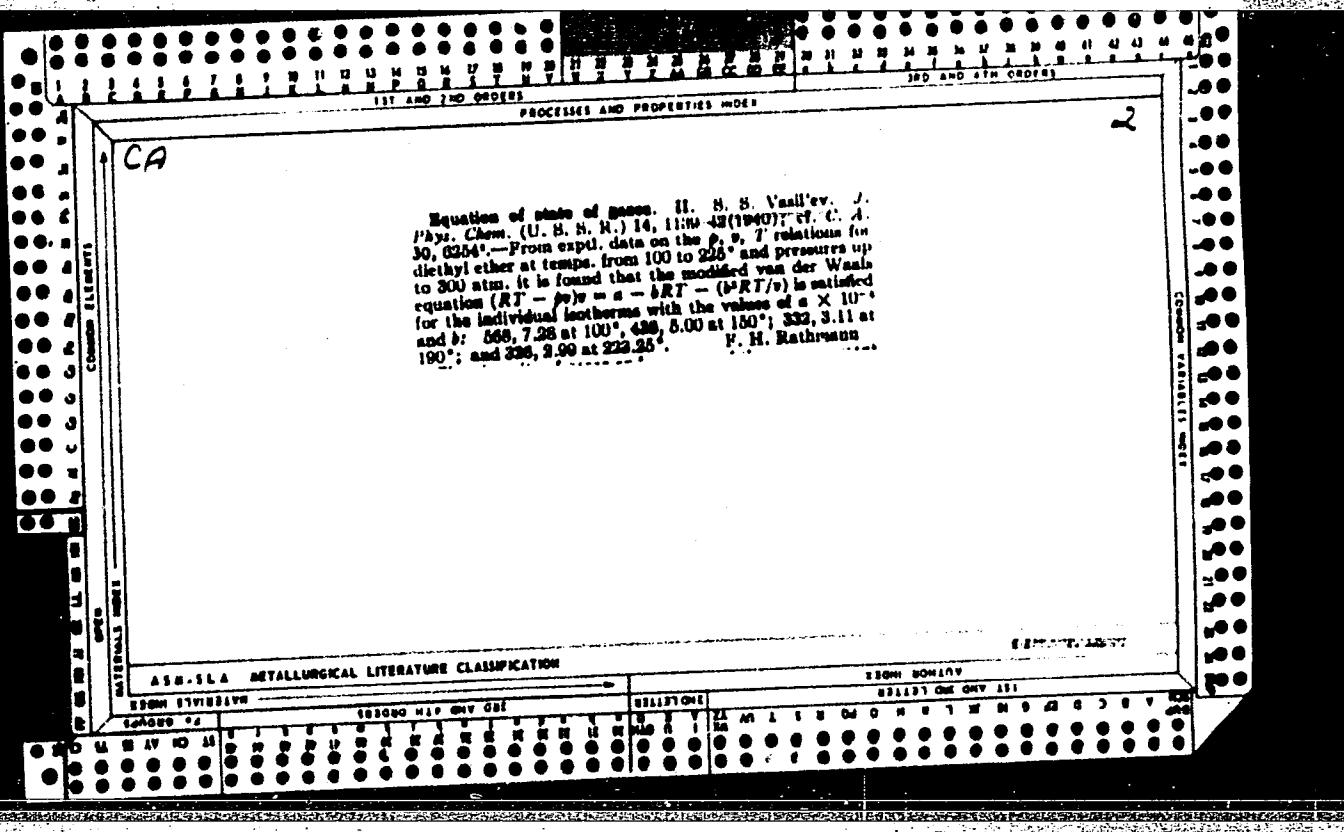


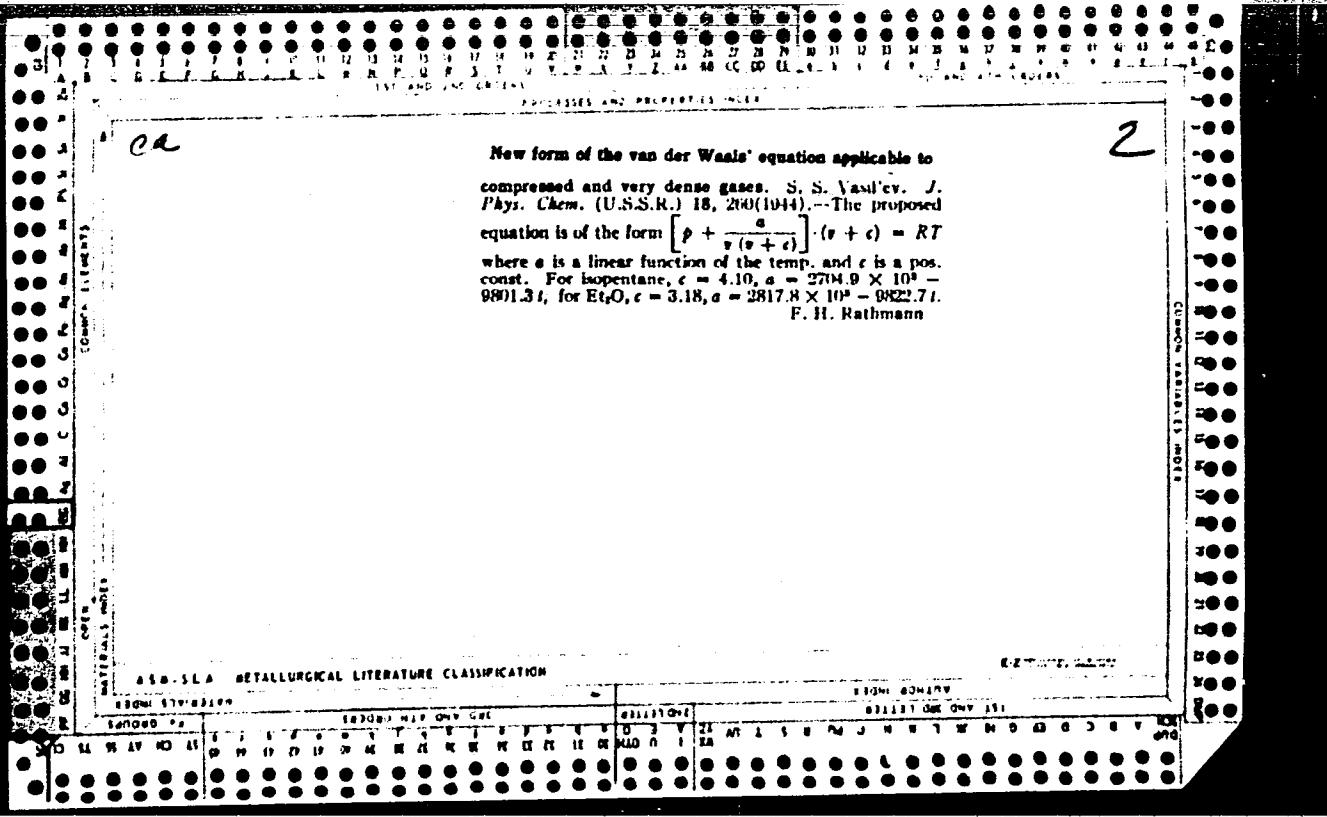


A correction to the article "Activation process during reactions in electric discharges," N. I. Kolobov, S. D. Vasil'ev and R. N. Iurmin, *J. Phys. Chem. U.S.S.R.* 11, 760 (1937); cf. *C. A.* 32, 6087, 8250. The reading of the spectrum appearing during the oxidation of N should be corrected to read: "To the emission energy of 1 e. v. corresponds the wave length 12,340 Å. The continuous spectrum found during the elec. oxidation of N is distributed at 200 mm. Hg into the region of 8000-9000 Å. wave lengths." After a further discussion K., V. and E. come to the conclusion that the difference between the full activation energy of N mole, calcd. from the wave lengths of the continuous spectrum, and the energy transferred during the collisions of "the second kind" can be regarded as the thermal energy of activation whose quantity lies in the limits from  $4.27 \pm 3.2 = 1.07$  e. v. = 21,680 cal./mol. to  $3.89 \pm 3.08 = 0.21$  e. v. = 4830 cal./mol.

W. R. Henn







VASIL'YEV, S. S.

PA 18T77

USSR/Chemistry - Hydrocarbon Oils  
Chemistry - Electrochemistry

Jun 1946

"A Kinetic Analysis of the Conversion of Methane to  
Acetylene and Hydrogen by Electrocracking," S. S.  
Vasil'yev, 22 pp Chair Physico-Chem., Lab. Catalysis &  
Gor'kogo Electro-Chem., Moscow State U.  
"Zhur Fiz Khim" Vol XX, No 6

Chemical formula and description of process used.  
Point is made that the experiments conducted enabled  
the observer to measure the output of the unsaturated  
compound under any given condition.

18T77

CA

3

Generalized formula for the calculation of effective electronic collisions in the electric discharge. S. S. Vasil'ev (Moscow State Univ.). *Vestn. Moskov. Univ.*, S, No. 3, Ser. Fiz.-Mat. i Estest. Nauk No. 3, 53-8(1950).—An Arrhenius-type formula is derived for the no.  $Z_{el}$  of effective collisions between electrons and atoms per unit length of the discharge,  $Z_{el} = Z \alpha_{el} \sigma_W e^{-\omega_1/kT_e}$ , where  $Z$  = total no. of "gas-kinetic" collisions per unit length of the discharge zone,  $\omega_1$  = excitation potential. The factor  $\alpha_{el}$  is a function of  $x = kT_e/\omega_1$  ( $\neq 1$ ), where  $\eta$  is defined by  $E_m = \eta \omega_1$ , i.e.  $\eta$  is the proportionality factor between  $\omega_1$  and the electronic energy at max. excitation;  $\eta = 2$  is only a special case. In the case of excitation of the higher electronic levels,  $x \rightarrow 0$ , and  $\alpha_{el} = |f_{el}/\omega_1|/(\eta - 1)$ , where  $f_{el}$  = excitation function; on the other hand, at very high  $T_e$ , the factor  $\alpha_{el} \rightarrow 0$ . The factor  $\sigma_W$  allows for the effect of chain and stepwise processes in the discharge zone. N. Tikhonov

CA

1 Energy balance and fixed energy distribution function in  
the physicochemical processes in electrical discharges  
S. S. Vasil'ev (Lomonosov State Univ., Moscow). *Zhur.  
Fiz. Khim.* 34, 1107-21 (1960).—An interpretative review.  
The analytical math. derivations verify previous work. 10  
references. Paul W. Howerton

CA

4

**Kinetic analysis of chemical processes in electric discharge.** S. S. Nasil'ev (Moscow State Univ.), *Vestn. Moskov. Univ. 6*, No. 8, Ser. Fiz.-Mat. i Estestven. Nauk No. 5, 79-90 (1951). - (1) With the contact time  $t$  of a chem. reaction in a flow through an elec. discharge, defined as the reciprocal  $\tau = 1/c$  of the feed rate  $c$  (cf. 6), the rate of formation of NO from  $N_2 + O_2$  is formulated  $dx/dt = k_1 - k_2 - k_3x$ , where  $x = \text{vol. fraction of NO in the outgoing gas}$ ,  $k_1$  is the zero-order rate const. of formation,  $k_2$  the 1st-order rate const. of decompr. of NO. The correctness of 1st-order kinetics of decompr. of NO in an elec. discharge, contrary to the 2nd order in thermal decompr., is borne out by agreement with exptl. data (see below). Without NO being added in advance, the integrated equation is  $x = (k_1/k_2)(1-e^{-k_2t})$ , with previous addn. of a vol. fraction  $x_0$  of NO,  $x = (k_1/k_2)(1 - e^{-k_2t}) + x_0e^{-k_2t}$ . At equil., at  $t \rightarrow \infty$ ,  $x = k_2/k_1 = x_0$ . The const.  $k_1$  can be detd. by varying  $t$  and measuring the corresponding  $x$ , by  $k_1 = r \ln(x_0/x - x_0)$ . One series of expts. with  $r$  varying from 0 to 135 l./hr. gave  $x$  varying from 5.8 to 1.5 vol. %, and  $k_1$  (av.) = 41 l./hr. While the values of  $k_1$  and  $k_2$  vary with the type and conditions of the elec. discharge, the validity of the above kinetic equations is confirmed by plotting  $x/x_0$  as a function of  $k_1 r$ , which gives one single curve,  $x/x_0 = 1 - e^{-k_2 t}$ , irrespective of the type of the discharge. A plot of the data of Shneerson and Vasil'ev (CA, 35, 3009), in the form of  $x$  as a function of  $t$ , shows a merger of the NO formation and decompn. curves into a common stationary  $x_0$ . Agreement between the exptl. points and the theoretical kinetic curves is obtained only with a 1st-order decompr. law for NO.

(2) Cracking of  $CH_4$  in elec. discharge can be described by the reactions:  $2CH_4 \rightarrow C_2H_2 + 3H_2$ ,  $2CH_4 \rightarrow C_2H_6 + 2H_2$ ,  $CH_4 \rightarrow C + 2H_2$ . The total cracking degree is defined by  $\Delta = x_1 - x_0$ , where  $x_1 = \text{vol. fraction of CH}_4 \text{ converted into unsatd. hydrocarbons}$ ;  $x_0 = 2q/(q + c)$ , where  $q = \text{vol. fraction of unsatd. hydrocarbons relative to the initial vol. of gas}$ . The yield of  $C_2H_2$  is  $\gamma = [C_2H_2]/\Delta$ . If, as usual,  $\gamma \approx 1$ , the vol. fraction  $x_1$  of unsatd. hydrocarbons in the outgoing gas is  $x_1 = x_0(1 - \Delta)$ . Experimentally, and in contrast to the 2nd order thermal cracking of  $CH_4$ , the cracking in an elec. discharge is 1st order. Hence,  $d\Delta/dt = k_1(\Delta - x_0) - k_2 x_1$ ,  $(k_1, k_2, k_3, k_4)$ , the subscripts refer to (2), (3), (4), (5).  $x_1 = x_0(1 - \Delta)$ ,  $(k_1, k_2, k_3, k_4)$ ,  $CH_4 \rightarrow C_2H_2 + H_2$ ,  $CH_4 \rightarrow C_2H_6 + H_2$ ,  $CH_4 \rightarrow C + 2H_2$ . Integration gives  $\Delta = \frac{1}{2}x_0e^{-k_1 t}$ . Finally  $\Delta = \frac{1}{2}x_0e^{-k_1 t}(1 - e^{-k_2 t})^{1/2}$ , an equation typical of consecutive reactions. On account of  $\Delta = 1 - e^{-k_1 t}$  and  $\gamma = \gamma_0$ , the final const. of unsatd. hydrocarbons is  $\alpha_1 = \frac{1}{2}\gamma_0e^{-k_1 t}(1 - e^{-k_2 t})^{1/2}[1 + \gamma_0(1 - e^{-k_2 t})]$ , i.e. passes through a max. as a function of  $t$ , this is confirmed by exptl. The max. of  $\alpha_1$  is the result of superposition of the increase of  $\Delta$  and the decrease of  $\sigma$  with time. The curves of  $\Delta$  as a function of  $k_1 r$ ,  $\alpha_1$  and of  $\sigma$  as a function of  $k_1 r$ , are independent of the type and the condition of the discharge. (3) Exptl. on cracking of  $CH_4$  in an elec. discharge zone, the rate const.  $k$  of any reaction in elec. discharge is proportional to the length  $l$  of the discharge zone,  $k = rl$ , where  $r$  is the rate const. per unit length. In a spherical discharge vessel, as long as its diam. is not too different from the diam. of the visible discharge zone, and at not too low a pressure,  $k_2$  for  $N_2 + O_2$  and  $k_1$  for  $CH_4$  are independent of the diam. of the vessel. This means that practically all the gas fed passes through the discharge zone. (4) In terms of the compn. of the gas, the "limiting energy" defined by  $\sigma = k_1/U$  (moles, kw, hr.), where  $U =$

Power input, passes through a max. just on the excess  $N_2$  side under low pressures (30 mm.), and on the excess  $N_2$  side under higher pressures (50 mm.). This shift of the max. rate of formation of  $N_2$  is characteristic of the reaction in diec. discharge; as in the thermal formation of  $N_2$  max. yield corresponds to a stoichiometric mix. In the case of the cracking of  $CH_4$ , previous claim, with  $H_2$  decreases the desorption rate const.,  $k_2$ , in such a way that the product  $k_1 k_2$  = initial concn. of  $CH_4$  in the  $CH_4 - H_2$  mix., remains approx. const. As cited, with the mean  $k_1 k_2$  in diec. discharge, with the expdt. 1. The constancy of  $k_1 k_2$  in electrocracking parallels the constancy of  $k_1 k_2$  in thermal cracking of  $CH_4$ . (6) As a function of the cond. of the diec. discharge, the rate const., per cm. length of the discharge zone,  $k_1 k_2$ , for the cracking of  $CH_4$ , increases rapidly and linearly with  $i$ . For the formation of  $N_2$ ,  $k_1 k_2$  increases with  $i$  much more slowly, and only up to some  $i_0$ , and then becomes very nearly independent of  $i$ . Empirically, an electrocracking of  $CH_4$  under 60-100 m. cond. is given by  $k_1 k_2 = 0.14 \cdot i^{0.1} + 0.14$  [W]  $- 0.14$  [W]<sup>2</sup>, where  $[W]$  is expressed in lux, term. of the power input, in w/cm.<sup>2</sup> and  $i$  in ma. The ratio  $k_1 k_2$  in  $CH_4/H_2$  mix., empirically,  $\sim 3.4$ . Some relations derived from the literature of unsteady-state reactions in the existing gas, when  $i$  is  $10^4$  amp., are given, and also cases of the energy expenditure of the discharge, in terms of hours. It increases with decreasing  $i$  or  $[W]$  attained to 2 kw hr. term. The thermal energy consumption being 1800 hr. term. for  $CH_4$ , the energy of electrocracking is about 1000 hr. term. for  $N_2$ . The formation rate const.,  $k_1$ , increases from 50 to 80 with the frequency increasing from 50 to 270 cps cycles/sec. Real  $[W] = 70$  w., pressure 180 mm., whereas the desorption rate const.,  $k_2$ , increases only from 10 to 16. Consequently, the increase of the sum of  $k_1 k_2$  is faster at higher frequency. The effect of variation of the  $N_2$  ratio is the same at first at low frequency. The higher frequency of high-frequency discharge leads only to a slight increase of the order of one-tenth to two times. These

SEREБRYAKOV, S.V., prof., doktor ekonom.nauk; GOGOL', B.I., dotsent;  
LIFITS, M.M., prof.; FEFILOV, A.I., dotsent; KISTANOV, Ya.A.,  
dotsent; GENKINA, L.S., dotsent; VASIL'YEV, S.S., dotsent;  
DNEPROVSKIY, S.P., prof.; PIROGOV, P.V., dotsent; SMOTRINA, N.A.,  
dotsent; KULIKOV, A.G., dotsent; KUZIN, N.I., dotsent; PISKUNOV, V.  
red. ; . . MUKHIN, Yu., tekhn.red.

[Economics of Soviet commerce] Ekonomika sovetskoi torgovli;  
uchebnoe posobie. Moskva, Gos.izd-vo polit.lit-ry, 1959. 478 p.

(MIRA 12:12)

(Russia--Commerce)

sics held in Tbilisi 14-22 Feb 1964/

SOURCE: AN SSSR, Izvestiya. Seriya fizicheskaya, v.29, no.1, 1965, 181-185

TOPIC TAGS: proton scattering, aluminum, nuclear spectroscopy, spin, parity, excited state.

Contd ... 4

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

SOURCE: Moskau, Universität, Versuch, Berlin, Physik, Astronomie, no. 1,

TOPIC NAME: Elektronenstrahlung, Elektronenstrahl, Elektronenstrahl, Elektronenstrahlung

ANALYST: [unclear]

\*\*\* L/C

ACCESSION NR: A9400000

scattering angle of 90° in the laboratory frame, in the energy interval 6.15--6.7 MeV. The experimental data were processed in the manner described in the earlier note and division of the different scattering channels indicating that the re-

ASSOCIATION: STYEF

SUBMITTED: CIAprc4

REF ID: A9400000

SUB CODE: NP

VASIL'YEV, S. S.

IA 248T13

"Kinetic Analysis of Chain Reactions: I. Kinetics of Chain Reactions Which Take Place With the Assistance of One Kind of Intermediate Active Particles," S. S. Vasil'yev, Inst of Light Industry imeni L. M. Kaganovich, Moscow

Zhur Fiz Khim, Vol 26, No 7, pp 1024-1035

This work aims at applying the methods of chem kinetics to an analysis of the general principles of chain reactions. It also strives to set up mathematical relationships characterizing these reactions, so as to compare results obtained by other (1)

researchers in this field. The author states that N. A. Shilov's eqs determine the fundamental form of mathematical relations which describe the kinetics of chain reactions. He criticizes N. N. Semenov for not recognizing the connection between Shilov's fundamental eq and the partial processes which determine the development of the chain reaction. Generalized kinetic consts made apparent the fundamental connection between the different values which can be applied for the characterization of chain reactions. The author applied the method for calcg the "Kinetic constant of diffusion" of active particles against the walls of a reaction container. This permitted a generalization of all (2)

248T13

cases of kinetics of chain reactions, and evaluation of the possible role which processes that take place at the walls of reaction containers will play in the creation of new active particles. (3)

248T13

1. VASIL'YEV, S. S.
2. USSR (600)
4. Chemical Reaction - Velocity
7. Analysis of kinetics of chain reactions. Part 2  
Kinetics of chain reactions taking place with the aid of several kinds  
of intermediate active particles.  
Zhur. fiz. khim. 26 no. 8, 1952
9. Monthly List of Russian Accessions, Library of Congress, January 1953. Unclassified.

VASIL'YEV, S. S.

IA 242T5

USSR/Chemistry - Electrochemistry

Nov 52

"Basic Elementary Processes and Physical Parameters Which Determine the Kinetics of Reactions in the Region of Electric Discharges," S. S. Vasil'yev, Moscow Technological Inst of Light Industry imeni L.M. Kaganovich.

"Zhur Fiz Khim" Vol 26, No 11, pp 1577-1590

The author offers a system of relationships unifying the various "basic" elementary processes which take place in the pos column of an elec discharge. Shows the connection between these processes and the basic phys parameters of the discharge, i.e. the current strength, the potential gradient, and the power of the discharge. Pays particular attention to the generalization of eqs of step-wise processes, whose significance is especially striking during great strengths of current. The author also shows the possibility of setting up a system of eqs permitting decision of the question regarding the concn of the excited mols in the region of discharge. This demonstrated to him the possibility of quantitatively connecting the rate of chem processes in a discharge with the elementary processes which take place in the region of the discharge and with the phys parameters of the discharge.

242T5

VASIL'EV, S.S.

✓ Corrections and remarks on "kinetic analysis of chain reactions." S. S. Vasil'ev, Zhur. Fiz. Khim. 27, 312 (1953); cf. C.A. 47, 6711f.—A no. of corrections are given along with a brief discussion of the integration of the kinetic equations that were developed. J. Rovtar Leach

check  
JRL  
PM

VASIL'YEV, S. S.

USSR/Chemistry - Reaction Kinetics

Jul 53

"Kinetic Analysis of Chain Reactions. III. General Integrals of Systems of Differential Kinetic Equations for the Initial Stages of Chain Reactions,"  
S. S. Vasil'yev, Technol Inst of Light Industry im  
L. M. Kaganovich, Moscow

Zhur Fiz Khim, Vol 27, No 7, pp 1081-1089

Calcd the general integrals of non-uniform systems of differential kinetic eqs for the initial stages of chain reactions. On the example of one of the possible cases of a chain reaction, carried out a complete soln of the general integrals and plotted

271T15

the corresponding curves. These curves indicate interesting characteristics of the development of chain reactions. Noted the possibility that a chain process may take place when concn of active particles fluctuates during the time that this concn increases.

VASIL'YEV, S. S.

USSR/Chemistry - Chain Reactions,  
Reaction Kinetics  
Sep 53

"Kinetic Analysis of Chain Reactions. IV. Distribution of Active Particles in Space and Calculation of Kinetic Constants of Diffusion," S. S. Vasil'yev, Technol Inst of Light Ind im L. M. Kaganovich, Moscow

Zhur Fiz Khim, Vol 27, No 9, pp 1410-1419

Calcd values of the integrals of an inhomogeneous system of kinetic differential eqs for an arbitrary distribn in space of the initial centers at which

269r27

chain reactions originate. Found that under certain conditions this distribn remains const in the vessel during the course of the chain reaction, so that calcn of the diffusion of active particles into the reaction zones becomes very simple. Calcd kinetic consts of diffusion of active particles for various shapes of the reaction vessel.

269r27

VASIL'YEV, S.S.

2  
0  
0  
0

*✓ The principal kinetic equations of chain reactions. S. S. Vasilev. Voprosy Khim. Kinetiki, Kataliza i Reaktsionnoi Speichnosti, Akad. Nauk S.S.R. 1955, 137-40.—A rigorous integration of systems of differential equations of chain reaction kinetics, in cases where the concn. changes of the reacting materials can be disregarded, was previously presented (C.A. 47, 5771; 49, 5935; preceding abstr.). The deduction of kinetic equations of chain reactions permits progressive increase in the concn. of active intermediate particles from the time of their generation until the drop in the concn. of the reacting substances can be disregarded. The kinetic curves of chain reactions may have entirely different forms at the initial moments, and these forms cannot be represented by any approximations. Special cases of general kinetic curves are discussed, and these cases may lead to the chain reaction kinetic equations formerly used. Differences between chain reactions and induced and conjugated reactions are shown not to be differences in principle.*

W. M. Sternberg

VASILYEV  
USSR/ Nuclear Physics

Card 1/2      Pub. 147 - 9/22

Authors : Vasiliyev, S. S.

Title : Kinetics of molecule excitation with electromagnetic and mechanical waves.  
Part 1. Importance of free energy cyclization during the excitation of  
molecules in structural units

Periodical : Zhur. fiz. khim. 29/11, 2018-2030, Nov 1955

Abstract : It is shown that the kinetics of molecule excitation processes with electro-  
magnetic waves in statistical and structural ensembles (concentration of  
molecules or atoms forms a structural ensemble), can be expressed by means  
of kinetic equations of a chain processes. It was found that a concentra-  
tion of excited molecules, in statistical and structural ensembles in a

Institution : Technological Institute of Light Industry im. L. M. Kaganovich, Moscow

Submitted : January 26, 1955

Card 2/2 APPROVED FOR RELEASE: 08/31/2001      CIA-RDP86-00513R001858910020-7"

Periodical : Zhur. fiz. khim. 29/11, 2018-2030, Nov 1955

Abstract : stationary state, can be represented by means of a very simple formula  
which includes the coefficients which take into consideration the effect  
of side energetic conversions in the molecules and macroscopic structural  
formations which concentrate or diffuse free radiation energy. Twelve  
references: 8 USSR, 3 Germ. and 1 USA (1933-1953).

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"

SHEYNIS, Ye.S., assistent; VASIL'YEV, S.S., prof.

Dielectric properties of dry and moist leather in the frequency range from 50 to 6,000 kc. Izv.vys.ucheb.zav.; tekhn.leg.prom. no.5:67-73 '58.  
(MIRA 12:2)

1. Moskovskiy tekhnologicheskiy institut legkoy promyshlennosti.  
(Leather--Electric properties)

AUTHORS:

Vasil'yev, S. S., Selivokhina, M. S. 76-32-6 17/46

TITLE:

The Kinetics of Electric Nitrogen Oxidation in the Auto-oscillating Discharge (Kinetika elektricheskogo okisleniya azota v avtokolebatel'nom razryade)

PERIODICALS:

Zhurnal fizicheskoy khimii 1958 Vol. 32, Nr 6, pp 1299-1303 (USSR)

ABSTRACT:

Electric autooscillations in the discharge zone of radiofrequency discharges with a frequency of one Mc were recently described by V. I. Granovskiy and L. N. Bykhovskiy (Refs 1-2). In the present paper high-frequency auto-oscillations were investigated occurring in the zone of electric low-frequency discharges in air at pressures ranging from a few mm of mercury column to atmospheric pressure. The experimental arrangement for the observation of the autooscillations, the discharge tube and the oscillographs obtained are described. In connection with the latter it is noted that at identical ignition conditions sometimes autooscillations may be seen on the oscillographs and sometimes not. Moreover changes in the color and the intensity

Card 1/3

The Kinetics of Electric Nitrogen Oxidation in the  
Autooscillating Discharge

76-32-6-77-4b

of the discharge spectrum can be observed at times. The evidence collected is assumed to give a proof for the earlier assumed theoretical conceptions that a possibility exists of the formation of actively bound oscillations of the electrons and optical mass elements. The results obtained and the experimental technique are described. It was observed among other things that a rise of the velocity and of the energy production of the process of electric oxidation of nitrogen corresponds to the formation of autooscillations. Detailed pertinent data are given. There are 9 figures, 1 table, and 7 references, 7 of which are Soviet.

ASSOCIATION: Tekhnologicheskiy institut iegkoy promyshlennosti, Moskva  
(Moscow, Technological Institute of Light Industry)

SUBMITTED: January 29, 1957

Card 2/3

The Kinetics of Electric Nitrogen Oxidation  
in the Autooscillating Discharge

76-52-6-11/46

1. Nitrogen--Oxidation    2. Electric discharges---Oscillation    3. Oscillations  
--Theory

Card 3/3

5 (4)

AUTHOR: Vasil'yev, S. S.

SOV/74-28-6-1/5

TITLE: Successes in the Investigation of Intermediate Reactions at the Photosynthetic Process of Carbohydrates (Uspekhi v issledovanii pomezhutochnykh reaktsiy protessa fotosintza uglevodov)

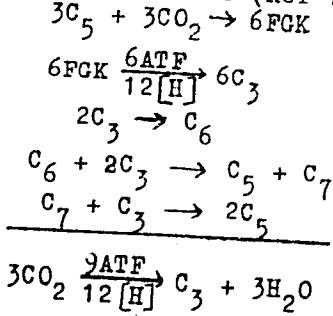
PERIODICAL: Uspekhi khimii, 1959, Vol 28, Nr 6, pp 647 - 668 (USSR)

ABSTRACT: The present paper tries to compile the most important results of investigation of intermediate reactions which together constitute the process of photosynthesis. The problem of photosynthesis can be divided into several part problems: 1) Mechanism of the "primary action" during which the absorption of light takes place and the absorbed energy is stabilized; 2) mechanism of processes which take place during the "obscure stage" of photosynthesis; 3) the amount of efficiency of light for the photosynthesis; 4) the particularities of absorption of light quantum by chlorophyll molecules. At present, the "obscure stage" is the problem worked out in the best way. These investigations are also of great importance for the explanation of the primary action, as they show the molecules

Card 1/4

Successes in the Investigation of Intermediate  
Reactions at the Photosynthetic Process of Carbohydrates SOV/74-28-6-1/5

forming during the primary action and being most important for the conversion of carbonic acid into carbohydrates. The mechanism of reactions taking place during the dark stage of the photosynthesis was primarily investigated on the basis of papers by Calvin and his collaborators (Refs 10-12). But also many other authors have been concerned with this problem (Refs 13-21). The reactions of the obscure stage can be represented by a somewhat simplified scheme (Ref 15):



Card 2/4

FGK - 3-phosphoglyceric acid, ATF - adenosintriphosphate.

Successes in the Investigation of Intermediate Reactions at the Photosynthetic Process of Carbohydrates      SOV/74-28-6-1/5

Many papers are dedicated to processes such as reduction, oxidation, phosphorylation, and hydrating, which are connected with the primary action (Refs 16, 18-20, 22-37). In the present survey, only a few of the papers on the mechanism of photosynthesis published in special publications are named. But they are of great importance for the introduction into the theory of intermediate reactions of photosynthesis. The above-mentioned part problems of the efficiency of the photosynthetic apparatus of green plants, as well as the analysis of particularities of the absorption of light by chlorophyll, are not considered. The number of papers on intermediate reactions of photosynthesis is steadily growing. The data published in these papers (Refs 38-50) are most contradictory. This is due to the circumstance that the process of photosynthesis is presumably a reaction which is closely connected with other biochemical processes in the organism of plants, and which can proceed in different ways. This shows that much work will have to be done to clarify the still unsolved prob-

Card 3/4

Successes in the Investigation of Intermediate  
Reactions at the Photosynthetic Process of Carbohydrates SCOV/74-28-6-1/5

lems of photosynthesis. There are 7 figures, 1 table, and 50  
references, 25 of which are Soviet.

ASSOCIATION: Moskovskiy tekhnologicheskiy in-t lekkoj promyshlennosti  
(Moscow Technological Institute of Light Industry)

Card 4/4

5 (4)

AUTHOR:

Vasil'yev, S. S. (Moscow)

COV/75-33-5-22/33

TITLE:

Kinetic Analysis of Chain Reactions (Kineticheskiy analiz tseptykh reaktsiy). 5. Transformation of the Form of Solutions of the Basic Equations for the Kinetics of Chain Reactions in the Case of Two Active Particles (5. Preobrazovaniye formy resheniy osnovnykh uravneniy kinetiki tseptykh reaktsiy dlya sluchaya dvukh aktivnykh chastits)

PERIODICAL:

Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 5,  
pp 1100-1110 (USSR)

ABSTRACT:

In previous papers (Refs 1-4) a fundamental differential equation for the kinetics mentioned in the title was investigated. The development of the chain reaction from the moment of the start ( $t = 0$ ) to a moment when the concentration decrease of the active particles may still be neglected is investigated in this paper. The coefficients  $s_1$ ,  $s_2$ ,  $r_{21}$ ,  $r_{12}$  were used ( $s_1$  and  $s_2$  = the general monomolecular constants for the reaction of the particles 1 and 2 in any direction,  $r_{12}$  = = the general kinetic molecular constant for the origin of

Card 1/3

Kinetic Analysis of Chain Reactions. 5. Transformation 30V/71-33-5-22/33  
of the Form of Solutions of the Basic Equations for the Kinetics of Chain  
Reactions in the Case of Two Active Particles

particle 1 by reaction of particle 2,  $r_{21}$  = the constant for  
the inverse course of the reaction. The ratio of the  
constants  $r_{ij}$  to  $s_i, s_j$  ( $i, j = 1, 2$ ) is expressed in

dimensionless quantities as  $\omega_{12} = \frac{r_{12}}{s_2}$ ;  $\omega_{21} = \frac{r_{21}}{s_1}$ ,  $\omega_{12}$  and  $\omega_{21}$

represent the mathematical expectation of the occurrence of  
the respective reaction. Their product  $\omega_{12}\omega_{21} = \omega$  is now  
introduced into the calculation. The kinetic equations obtained  
by determinant computation permit a direct determination of  
the effect of the formation rates, the initial concentration,  
and the reaction coefficients of the active particles on the  
temporal variation of their concentration. The cases  
 $\omega < 1$ ,  $\omega > 1$ , and  $\omega \rightarrow 1$  are analyzed. With  $\omega < 1$  the  
concentration of the active particles reaches stationary state  
(Formula 54), with  $\omega > 1$  the linear concentration increase of  
the active particle is substituted by an exponential one

Card 2/3

Kinetic Analysis of Chain Reactions.

SOV/76-33-5-22/33

5. Transformation of the Form of Solutions of the Basic Equations for the Kinetics of Chain Reactions in the Case of Two Active Particles

(Formula 56). With  $\omega \rightarrow 1$  the time of the occurrence of the states (54) and (56) becomes infinite. Thus  $\omega \rightarrow 1$  separates the range of the processes leading to stationary states from those proceeding with self-acceleration and may be considered the definition of the "lower limit" of chain reactions. With regard to separation of chain reactions into reactions with ramification and reactions without ramification it is stated that stationary states can also occur in the case of reactions with ramification if, e. g.  $\omega_{12} > 1$ , but  $\omega_{21} \ll 1$ , so that  $\omega$  remains  $< 1$ . There are 8 references, 7 of which are Soviet.

ASSOCIATION: Tekhnologicheskiy institut legkoy promyshlennosti Moskva  
(Technological Institute of Light Industry, Moscow)

SUBMITTED: October 25, 1957

Card 3/3

VASIL'YEV, S.S.; KOMAROV, V.V.; POPOVA, A.M.

Decay reactions of carbon and oxygen nuclei induced by 15-29 Mev.  
protons. Zhur. eksp. i teor. fiz. 43 no.3:737-748 '62. (MIRA 15:10)

1. Institut yadernoy fiziki Moskovskogo gosudarstvennogo universiteta.  
(Nuclear reactions) (Protons)

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7

CHALMERS, T.M., Naval Research Laboratory, Washington, D.C.  
Technical Report, 1951.

Kinetics of the evaporation of ammonium nitrate from a  
solution of ammonium nitrate. Naval Research Laboratory, 1951  
U.S.

AMMONIUM NITRATE

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001858910020-7"